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(54) Title: **NOVEL ARYLALKANE-SULFONAMIDES**

(57) Abstract: The invention relates to novel aryl-alkane-sulfonamides and their use as active ingredients in the preparation of pharmaceutical compositions. The invention also concerns related aspects including processes for the preparation of the compounds, pharmaceutical compositions containing one or more of those compounds and especially their use as endothelin receptor antagonists.

**WO 02/24665 A1**

## NOVEL ARYLALKANE-SULFONAMIDES

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The present invention relates to novel arylalkane-sulfonamides of the general formula I and their use as active ingredients in the preparation of pharmaceutical compositions. The invention also concerns related aspects including processes for the preparation of the compounds, pharmaceutical compositions containing  
10 one or more compounds of the general formula I and especially their use as endothelin receptor antagonists.

Endothelins (ET-1, ET-2, and ET-3) are 21-amino acid peptides produced and active in almost all tissues (Yanagisawa M et al.: Nature (1988) 332:411).  
15 Endothelins are potent vasoconstrictors and important mediators of cardiac, renal, endocrine and immune functions (McMillen MA et al.: J Am Coll Surg (1995) 180:621). They participate in bronchoconstriction and regulate neurotransmitter release, activation of inflammatory cells, fibrosis, cell proliferation and cell differentiation (Rubanyi GM et al.: Pharmacol Rev (1994)  
20 46:328).

Two endothelin receptors have been cloned and characterized in mammals (ET<sub>A</sub>, ET<sub>B</sub>) (Arai H et al.: Nature (1990) 348:730; Sakurai T et al.: Nature (1990) 348:732). The ET<sub>A</sub> receptor is characterized by higher affinity for ET-1 and ET-2  
25 than for ET-3. It is predominant in vascular smooth muscle cells and mediates vasoconstricting and proliferative responses (Ohlstein EH et al.: Drug Dev Res (1993) 29:108). In contrast, the ET<sub>B</sub> receptor has equivalent affinity for the 3 endothelin isopeptides and binds the linear form of endothelin, tetra-ala-endothelin, and sarafotoxin S6C (Ogawa Y et al.: BBRC (1991) 178:248). This  
30 receptor is located in the vascular endothelium and smooth muscles, and is also particularly abundant in lung and brain. The ET<sub>B</sub> receptor from endothelial cells mediates transient vasodilator responses to ET-1 and ET-3 through the release of nitric oxide and/or prostacyclin whereas the ET<sub>B</sub> receptor from smooth muscle

cells exerts vasoconstricting actions (Sumner MJ et al.: Brit J Pharmacol (1992) 107:858). ET<sub>A</sub> and ET<sub>B</sub> receptors are highly similar in structure and belong to the superfamily of G-protein coupled receptors.

- 5 A pathophysiological role has been suggested for ET-1 in view of its increased plasma and tissue levels in several disease states such as hypertension, sepsis, atherosclerosis, acute myocardial infarction, congestive heart failure, renal failure, migraine and asthma. As a consequence, endothelin receptor antagonists have been studied extensively as potential therapeutic agents. Endothelin
- 10 receptor antagonists have demonstrated preclinical and/or clinical efficacy in various diseases such as cerebral vasospasm following subarachnoid hemorrhage, heart failure, pulmonary and systemic hypertension, neurogenic inflammation, renal failure and myocardial infarction.
- 15 Today, no endothelin receptor antagonist is marketed yet, several are in clinical trials. However, these molecules possess a number of weaknesses such as complex synthesis, low solubility, high molecular weight, poor pharmacokinetics, or safety problems (e.g. liver enzyme increases).
- 20 The inhibitory activity of the compounds of general formula I on endothelin receptors can be demonstrated using the test procedures described hereinafter.

*For the evaluation of the potency and efficacy of the compounds of the general formula I the following tests were used:*

5    **1) Inhibition of endothelin binding to membranes from CHO cells carrying human ET receptors:**

For competition binding studies, membranes of CHO cells expressing human recombinant ET<sub>A</sub> or ET<sub>B</sub> receptors were used. Microsomal membranes from recombinant CHO cells were prepared and the binding assay made as previously  
10    described (Breu V., et al, FEBS Lett 1993; 334:210).

The assay was performed in 200 uL 50 mM Tris/HCl buffer, pH 7.4, including 25 mM MnCl<sub>2</sub>, 1 mM EDTA and 0.5% (w/v) BSA in polypropylene microtiter plates. Membranes containing 0.5 ug protein were incubated for 2 h at 20°C with 8 pM  
15    [<sup>125</sup>I]ET-1 (4000 cpm) and increasing concentrations of unlabelled antagonists. Maximum and minimum binding were estimated in samples without and with 100 nM ET-1, respectively. After 2 h, the membranes were filtered on filterplates containing GF/C filters (Unifilterplates from Canberra Packard S.A. Zürich, Switzerland). To each well, 50 uL of scintillation cocktail was added (MicroScint  
20    20, Canberra Packard S.A. Zürich, Switzerland) and the filter plates counted in a microplate counter (TopCount, Canberra Packard S.A. Zürich, Switzerland).

All the test compounds were dissolved, diluted and added in DMSO. The assay was run in the presence of 2.5% DMSO which was found not to interfere  
25    significantly with the binding. IC<sub>50</sub> was calculated as the concentration of antagonist inhibiting 50 % of the specific binding of ET-1. For reference compounds, the following IC<sub>50</sub> values were found: ET<sub>A</sub> cells: 0.075 nM (n=8) for ET-1 and 118 nM (n=8) for ET-3; ET<sub>B</sub> cells: 0.067 nM (n=8) for ET-1 and 0.092 nM (n=3) for ET-3.

30

The IC<sub>50</sub> values obtained with compounds of general formula I are given in Table 1.

**Table 1:**

Compound of Example	IC <sub>50</sub> ET <sub>A</sub> [nM]	IC <sub>50</sub> ET <sub>B</sub> [nM]
Example 1	27	6650
Example 2	20	899
Example 5	3	323
Example 7	4	3310
Example 10	9	2410
Example 13	4	3680
Example 14	6	2230
Example 19	3	1930
Example 29	10	406
Example 39	3	261
Example 46	7	2360
Example 47	24	2720
Example 49	4	2490
Example 52	5	1770
Example 61	10	1140
Example 71	115	>10000
Example 81	24	824

**2) Inhibition of endothelin-induced contractions on isolated rat aortic rings (ET<sub>A</sub> receptors) and rat tracheal rings (ET<sub>B</sub> receptors):**

The functional inhibitory potency of the endothelin antagonists was assessed by their inhibition of the contraction induced by endothelin-1 on rat aortic rings (ET<sub>A</sub> receptors) and of the contraction induced by sarafotoxin S6c on rat tracheal rings (ET<sub>B</sub> receptors). Adult Wistar rats were anesthetized and exsanguinated. The thoracic aorta or trachea were excised, dissected and cut in 3-5 mm rings. The endothelium/epithelium was removed by gentle rubbing of the intimal surface. Each ring was suspended in a 10 ml isolated organ bath filled with Krebs-

Henseleit solution (in mM; NaCl 115, KCl 4.7, MgSO<sub>4</sub> 1.2, KH<sub>2</sub>PO<sub>4</sub> 1.5, NaHCO<sub>3</sub> 25, CaCl<sub>2</sub> 2.5, glucose 10) kept at 37°C and gassed with 95% O<sub>2</sub> and 5% CO<sub>2</sub>. The rings were connected to force transducers and isometric tension was recorded (EMKA Technologies SA, Paris, France). The rings were stretched to a resting tension of 3 g (aorta) or 2 g (trachea). Cumulative doses of ET-1 (aorta) or sarafotoxin S6c (trachea) were added after a 10 min incubation with the test compound or its vehicle. The functional inhibitory potency of the test compound was assessed by calculating the concentration ratio, i.e. the shift to the right of the EC<sub>50</sub> induced by different concentrations of test compound. EC<sub>50</sub> is the concentration of endothelin needed to get a half-maximal contraction, pA<sub>2</sub> is the negative logarithm of the antagonist concentration which induces a two-fold shift in the EC<sub>50</sub> value.

The pA<sub>2</sub> values obtained with compounds of formula I are given in Table 2.

**Table 2:**

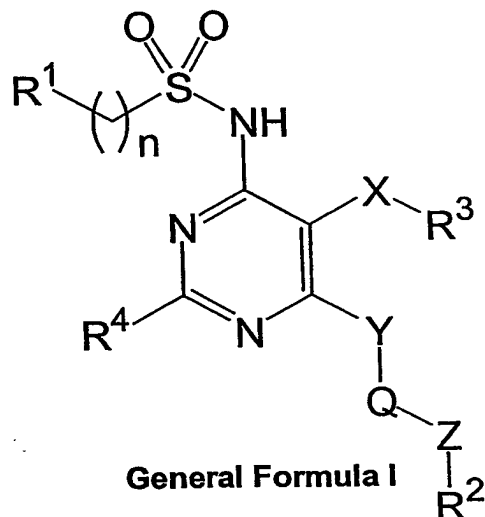
Compound of Example	pA <sub>2</sub> (aortic rings)	pA <sub>2</sub> (trachea)
Example 5	8.38	7.02
Example 7	8.83	7.07
Example 8	7.43	-
Example 34	7.67	-
Exempl 61	7.83	7.07
Example 75	7.76	-

Because of their ability to inhibit the endothelin binding, the described compounds can be used for treatment of diseases which are associated with an increase in vasoconstriction, proliferation or inflammation due to endothelin. Examples of such diseases are hypertension, coronary diseases, cardiac insufficiency, renal and myocardial ischemia, renal failure, cerebral ischemia, dementia, migraine, subarachnoidal hemorrhage, Raynaud's syndrome, portal hypertension and pulmonary hypertension. They can also be used for

- atherosclerosis, prevention of restenosis after balloon or stent angioplasty, inflammation, stomach and duodenal ulcer, cancer, prostatic hypertrophy, erectile dysfunction, hearing loss, amaurosis, chronic bronchitis, asthma, gram negative septicemia, shock, sickle cell anemia, glomerulonephritis, renal colic,
- 5 glaucoma, therapy and prophylaxis of diabetic complications, complications of vascular or cardiac surgery or after organ transplantation, complications of cyclosporin treatment, pain as well as other diseases presently known to be related to endothelin.
- 10 The compounds can be administered orally, rectally, parenterally, e.g. by intravenous, intramuscular, subcutaneous, intrathecal or transdermal administration or sublingually or as ophthalmic preparation or administered as aerosol. Examples of applications are capsules, tablets, orally administered suspensions or solutions, suppositories, injections, eye-drops, ointments or
- 15 aerosols/nebulizers.

- Preferred applications are intravenous, intramuscular, or oral administrations as well as eye drops. The dosage used depends upon the type of the specific active ingredient, the age and the requirements of the patient and the kind of
- 20 application. Generally, dosages of 0.1 – 50 mg / kg body weight per day are considered. The preparations with compounds can contain inert or as well pharmacodynamically active excipients. Tablets or granules, for example, could contain a number of binding agents, filling excipients, carrier substances or diluents.

The present invention relates to arylethene-sulfonamides of the general formula I,



5

wherein

$R^1$  and  $R^2$  represent aryl; heteroaryl;

$R^3$  represents phenyl; mono-, di- or tri-substituted phenyl substituted with lower alkyl, lower alkenyl, lower alkynyl, phenyl, lower alkoxy, amino, lower alkylamino, amino-lower alkyl, trifluoromethyl, trifluoromethoxy, halogen, lower alkylthio, hydroxy, hydroxy-lower alkyl, cyano, carboxyl, lower alkanoyl, formyl; benzofuranyl; aryl; heteroaryl;

$R^4$  represents hydrogen; halogen; trifluoromethyl; lower alkyl; lower alkyl-amino; lower alkoxy; lower alkyl-sulfonyl; lower alkyl-sulfinyl; lower alkylthio; lower alkylthio-lower alkyl; hydroxy-lower alkyl; lower alkyl-oxy-lower alkyl; hydroxy-lower alkyl-oxy-lower alkyl; hydroxy-lower alkyl-amino; lower alkyl-amino-lower alkyl; amino; di-lower alkyl-amino; [N-(hydroxy-lower alkyl)-N-(lower alkyl)]-amino; aryl; aryl-amino; aryl-lower alkyl-amino; aryl-thio; aryl-lower alkyl-thio; aryloxy; aryl-lower alkyl-oxy; aryl-lower alkyl; aryl-sulfinyl; heteroaryl; heteroaryl-oxy; heteroaryl-lower alkyl-oxy; heteroaryl-amino; heteroaryl-lower alkyl-amino; heteroaryl-thio; heteroaryl-lower alkyl-thio; heteroaryl-lower alkyl; heteroaryl-sulfinyl; heterocyclyl; heterocyclyl-lower alkyl-oxy; heterocyclyl-oxy; heterocyclyl-



amino; heterocyclyl-lower alkyl-amino; heterocyclyl-thio; heterocyclyl-lower alkyl-thio; heterocyclyl-lower alkyl; heterocyclyl-sulfinyl; cycloalkyl; cycloalkyl-oxy; cycloalkyl-lower alkyl-oxy; cycloalkyl-amino; cycloalkyl-lower alkyl-amino; cycloalkyl-thio; cycloalkyl-lower alkyl-thio; cycloalkyl-lower alkyl; cycloalkyl-sulfinyl;

5

X represents oxygen; sulfur; NH; CH<sub>2</sub> or a bond;

10 Y represents oxygen; sulfur or -NH-;

Z represents oxygen; sulfur, -NH- or a bond;

Q represents -(CH<sub>2</sub>)<sub>k</sub>-; -(CH<sub>2</sub>)<sub>m</sub>-C≡C-(CH<sub>2</sub>)<sub>p</sub>-, in case p represents 0 (zero), Z represents a bond; -CH<sub>2</sub>-cyclopropyl-CH<sub>2</sub>-;

15

k represents the numbers 2, 3, 4, 5, or 6;

m represents the numbers 1, 2, or 3;

20

p represents the numbers 0, 1, 2 or 3;

n represents the numbers 1, 2, or 3;

25 and pure diastereomers, mixtures of diastereomers, diastereomeric racemates, mixtures of diastereomeric racemates and the meso-forms and pharmaceutically acceptable salts thereof.

In the definitions of the general formula I – if not otherwise stated – the expression lower alkyl or lower alkoxy means straight and branched chain groups with one to seven carbon atoms, preferably 1 to 4 carbon atoms. Examples of lower alkyl and lower alkoxy groups are methyl, ethyl, n-propyl, isopropyl, n-butyl, isobutyl, sec.-butyl, tert.-butyl, pentyl, hexyl, heptyl, methoxy,

30

ethoxy, propoxy, n-butoxy, iso-butoxy, sec.-butoxy and tert.-butoxy. Lower alkylendioxy-groups are preferably methylene-dioxy, ethylene-dioxy, propylen-dioxy and butylen-dioxy groups. Examples of lower alkanoyl-groups are acetyl, propanoyl and butanoyl. Lower alkenylen means e.g.vinylen, propenylen and butenylen. Lower alkenyl and lower alkynyl means groups like ethylene, propylen, butylen, 2-methyl-propenyl, and ethinylen, propinylen, butinylen, pentinylen, 2-methyl-pentinylen etc. Lower alkenyloxy means allyloxy, vinyloxy, propenyloxy and the like. The expression **cycloalkyl** means a saturated cyclic hydrocarbon ring with 3 to 7 carbon atoms , e.g. cyclopropyl, cyclobutyl, cyclopentyl, cyclohexyl and cycloheptyl, which may be substituted with lower alkyl, hydroxy-lower alkyl, amino-lower alkyl, lower alkoxy-lower alkyl and lower alkenylen groups. The expression **heterocyclyl** means saturated or partially unsaturated four, five-, six- or seven-membered rings containing one or two nitrogen, oxygen or sulfur atoms which may be the same or different and which rings may be adequately substituted with lower alkyl, amino, nitro, hydroxy, lower alkoxy, e.g. piperidinyl, morpholinyl, thiomorpholinyl, piperazinyl, tetrahydropyranyl, dihydropyranyl, 1,4-dioxanyl, pyrrolidinyl, tetrahydrofuranyl, dihydropyrrolyl, dihydroimidazolyl, dihydropyrazolyl, pyrazolidinyl, 5-oxo-1,2,4-oxadiazolyl, 5-oxo-1,2,4-thiadiazolyl, 5-thioxo-1,2,4-oxadiazolyl, 2-oxo-1,2,3,5-oxathiadiazolyl etc. (e.g.[7]) and substituted derivatives of such rings with substituents as outlined above. The expression **heteroaryl** means six-membered aromatic rings containing one to four nitrogen atoms, benzofused six-membered aromatic rings containing one to three nitrogen atoms, five-membered aromatic rings containing one oxygen or one nitrogen or one sulfur atom, benzo- fused five-membered aromatic rings containing one oxygen or one nitrogen or one sulfur atom, five membered aromatic rings containig an oxygen and nitrogen atom and benzo fused derivatives thereof, five membred aromatic rings containing a sulfur and a nitrogen atom and benzo fused derivatives thereof, five-membered aromatic rings containing two nitrogen atoms and benzo fused derivatives thereof, five membered aromatic rings containing three nitrogen atoms and benzo fused derivatives thereof or the tetrazolyl ring; e.g. furanyl, thienyl, pyrrolyl, pyridinyl, pyrimidinyl, indolyl, quinolinyl, isoquinolinyl, imidazolyl, triazinyl, thiazinyl, thiazolyl, isothiazolyl, pyridazinyl, oxazolyl, isoxazolyl, etc.

whereby such rings may be substituted with lower alkyl, lower alkenyl, amino, amino-lower alkyl, halogen, hydroxy, lower alkoxy, trifluoromethoxy, trifluoromethyl, carboxyl, carboxamidyl, thioamidyl, amidinyl, lower alkoxy, cyano, hydroxy-lower alkyl, lower alkoxy-lower alkyl or another heteroaryl- (preferably  
5 tetrazolyl) or heterocyclyl-ring (preferably 5-oxo-1,2,4-oxadiazolyl, 5-oxo-1,2,4-triazolyl, 5-oxo-1,2,4-thiadiazolyl, 5-thioxo-1,2,4-oxadiazolyl or 2-oxo-1,2,3,5-oxathiadiazolyl (e.g. [7])). The expression *aryl* represents unsubstituted as well as mono-, di- or tri-substituted aromatic rings with 6 to 10 carbon atoms like phenyl or naphthyl rings which may be substituted with aryl, halogen, hydroxy,  
10 lower alkyl, lower alkenyl, lower alkynyl, lower alkoxy, lower alkenyloxy, lower alkynyl-lower alkyl-oxy, lower alkenylen, lower alkyleneoxy, lower alkyleneoxy or lower alkylendioxy forming with the phenyl ring a five- or six-membered ring, hydroxy-lower alkyl, hydroxy-lower alkenyl, hydroxy-lower alkyl-lower alkynyl, lower alkoxy-lower alkyl, lower alkoxy-lower alkoxy, trifluoromethyl,  
15 trifluoromethoxy, cycloalkyl, hydroxy-cycloalkyl, heterocyclyl, heteroaryl.

It is understood that the substituents outlined relative to the expressions cycloalkyl, heterocyclyl, heteroraryl and aryl have been omitted in the definitions of the general formulae I to V in claims 1 to 11 for clarity reasons but the  
20 definitions in formulae I to V and in claims 1 to 11 should be read as if they are included therein.

Especially preferred compounds are compounds of general formula I wherein  $R^3$  represents phenyl or mono-substituted phenyl substituted with lower alkoxy, especially methoxy and X represents oxygen.  
25

A second group of especially preferred compounds of general formula I are the compounds wherein  $R^3$  represents phenyl or mono-substituted phenyl substituted with lower alkoxy, especially methoxy, and wherein X, Y and Z  
30 represent oxygen.

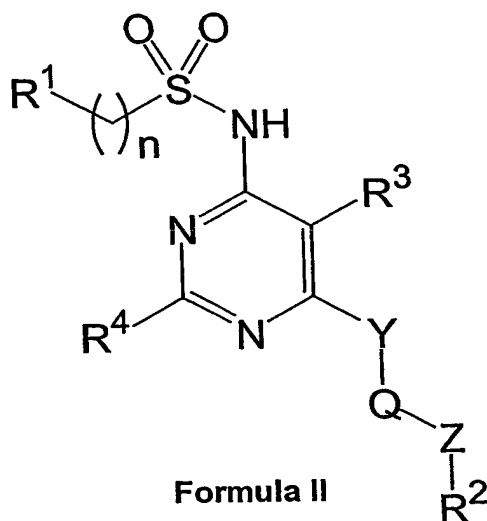
A third group of especially preferred compounds of general formula I are the compounds wherein  $R^3$  represents phenyl or mono-substituted phenyl

substituted with lower alkoxy, especially methoxy, and wherein X, Y and Z represent oxygen and Q represents  $-(CH_2)_k-$  with  $k = 2$  or  $3$ .

5 A fourth group of especially preferred compounds of general formula I are the compounds wherein  $R^2$  represents heteroaryl,  $R^3$  represents phenyl or mono-substituted phenyl substituted with lower alkoxy, especially methoxy, X, Y and Z represent oxygen and Q represents  $-(CH_2)_k-$  with  $k = 2$  or  $3$ .

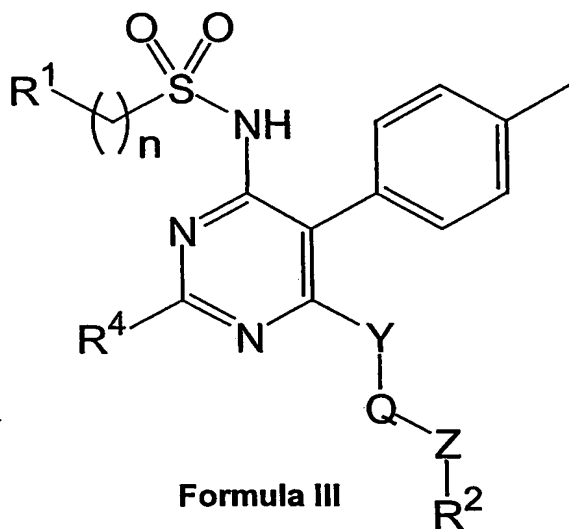
10 A fifth group of especially preferred compounds of general formula I are the compounds wherein  $R^2$  represents heteroaryl,  $R^3$  represents phenyl or mono-substituted phenyl substituted with halogen, lower alkyl, lower alkenyl, methoxy, amino, lower alkyl-amino, lower alkyl-thio, hydroxy, hydroxymethyl and lower alkanoyl; X, Y and Z represent oxygen, Q represents  $-(CH_2)_2-$ .

15 Another group of preferred compounds are compounds of formula II



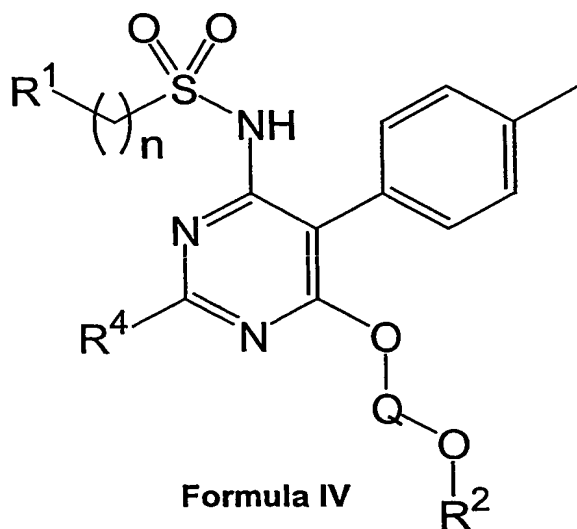
20 wherein  $R^1$ ,  $R^2$ ,  $R^3$ ,  $R^4$ , Y, Q, Z, and n are as defined in general formula I above, and pharmaceutically acceptable salts of compounds of formula II.

Also preferred are compounds of formula III



- 5 wherein  $R^1$ ,  $R^2$ ,  $R^4$ ,  $Y$ ,  $Q$  and  $Z$  and  $n$  are as defined in general formula I above, and pharmaceutically acceptable salts of compounds of formula III.

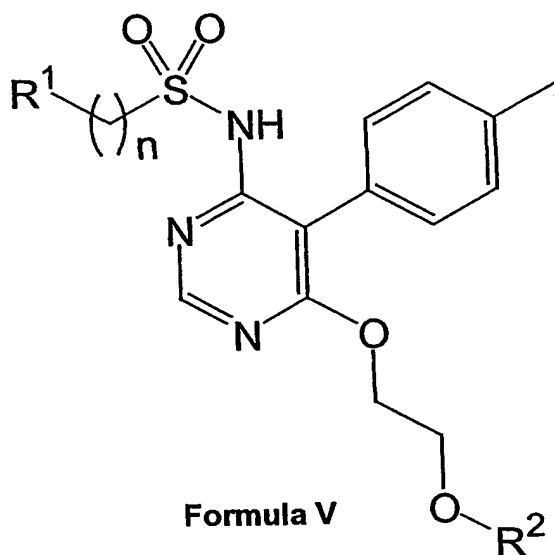
Also preferred are compounds of formula IV



10

wherein  $R^1$ ,  $R^2$ ,  $R^4$ ,  $Q$ , and  $n$  are as defined in general formula I above, and pharmaceutically acceptable salts of compounds of formula IV.

Another especially preferred group of compounds are compounds of formula V



- wherein  $R^1$  and  $R^2$  are as defined in general formula I above,  
 5 and pharmaceutically acceptable salts thereof.

Especially preferred compounds among the group of compounds of formula V are those wherein  $R^2$  represents heteroaryl.

- 10 The expression pharmaceutically acceptable salts encompasses either salts with inorganic acids or organic acids like hydrohalogenic acids, e.g. hydrochloric or hydrobromic acid; sulfuric acid, phosphoric acid, nitric acid, citric acid, formic acid, acetic acid, maleic acid, tartaric acid, methylsulfonic acid, p- toluolsulfonic acid and the like or in case the compound of formula I is acidic in nature with an  
 15 inorganic base like an alkali or earth alkali base, e.g. sodium hydroxide, potassium hydroxide, calcium hydroxide etc.

- The compounds of the general formula I might have one or more asymmetric carbon atoms and may be prepared in form of optically pure enantiomers or diastereomers, mixtures of enantiomers or diastereomers, diastereomeric  
 20 racemates, mixtures of diastereomeric racemates and also in the meso-form. The present invention encompasses all these forms. Mixtures may be separated in a manner known per se, i.e. by column chromatography, thin layer chromatography, HPLC, crystallization etc.

Because of their ability to inhibit the endothelin binding, the described compounds of the general formula I and their pharmaceutically acceptable salts may be used for treatment of diseases which are associated with an increase in vasoconstriction, proliferation or inflammation due to endothelin. Examples of such diseases are hypertension, coronary diseases, cardiac insufficiency, renal and myocardial ischemia, renal failure, cerebral ischemia, dementia, migraine, subarachnoidal hemorrhage, Raynaud's syndrome, portal hypertension and pulmonary hypertension. They can also be used for atherosclerosis, prevention of restenosis after balloon or stent angioplasty, inflammation, stomach and duodenal ulcer, cancer, prostatic hypertrophy, erectile dysfunction, hearing loss, amaurosis, chronic bronchitis, asthma, gram negative septicemia, shock, sickle cell anemia, glomerulonephritis, renal colic, glaucoma, therapy and prophylaxis of diabetic complications, complications of vascular or cardiac surgery or after organ transplantation, complications of cyclosporin treatment, pain, as well as other diseases presently known to be related to endothelin.

These compositions may be administered in enteral or oral form e.g. as tablets, dragees, gelatine capsules, emulsions, solutions or suspensions, in nasal form like sprays or rectically in form of suppositories. These compounds may also be administered in intramuscular, parenteral or intravenous form, e.g. in form of injectable solutions.

These pharmaceutical compositions may contain the compounds of formula I as well as their pharmaceutically acceptable salts in combination with inorganic and/or organic excipients which are usual in the pharmaceutical industry like lactose, maize or derivatives thereof, talcum, stearinic acid or salts of these materials.

For gelatine capsules vegetable oils, waxes, fats, liquid or half-liquid polyols etc. may be used. For the preparation of solutions and sirups e.g. water, polyols, saccharose, glucose etc. are used. Injectables are prepared by using e.g. water, polyols, alcohols, glycerin, vegetable oils, lecithin, liposomes etc. Suppositories

are prepared by using natural or hydrogenated oils, waxes, fatty acids (fats ), liquid or half-liquid polyols etc.

5 The compositions may contain in addition preservatives, stabilisation improving substances, viscosity improving or regulating substances, solubility improving substances, sweeteners, dyes, taste improving compounds, salts to change the osmotic pressure, buffer, anti-oxidants etc.

10 The compounds of formula I may also be used in combination with one or more other therapeutically useful substances e.g.  $\alpha$ - and  $\beta$ -blockers like phentolamine, phenoxybenzamine, atenolol, propranolol, timolol, metoprolol, carteolol etc.; Vasodilators like hydralazine, minoxidil, diazoxide, flosequinan etc.; Calcium-antagonists like diltiazem, nicardipine, nimodipine, verapamil, nifedipine etc.; ACE-inhibitors like cilazapril, captopril, enalapril, lisinopril etc.; Potassium  
15 activators like pinacidil etc.; Angiotensin II antagonists; Diuretics like hydrochlorothiazide, chlorothiazide, acetolamide, bumetanide, furosemide, metolazone, chlortalidone etc.; Sympatholitics like methyldopa, clonidine, guanabenz, reserpine etc.; and other therapeutics which serve to treat high blood pressure or any cardiac disorders.

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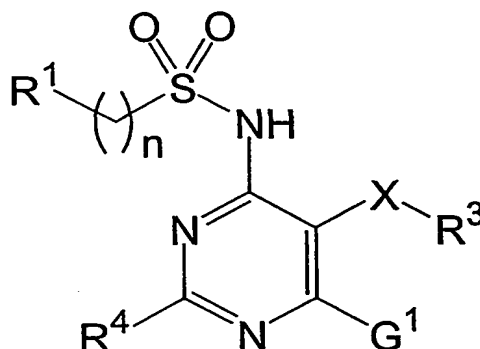
The dosage may vary within wide limits but should be adapted to the specific situation. In general the dosage given daily in oral form should be between about 3 mg and about 3 g, preferably between about 10 mg and about 1 g, especially preferred between 5 mg and 300 mg, per adult with a body weight of about 70  
25 kg. The dosage should be administered preferably in 1 to 3 doses per day which are of equal weight. As usual children should receive lower doses which are adapted to body weight and age.



Compounds of the **general formula I** of the present invention can be prepared according to the general sequence of reactions outlined below. For simplicity and clarity reasons sometimes only parts of the synthetic possibilities which lead to compounds of general formula I are described. The literature references given in  
 5 brackets [ ] are set forth at the end of this paragraph.

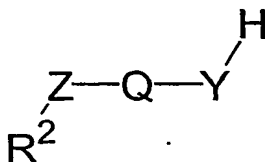
**Possibility A:**

The desired compounds of general formula I can be prepared by reacting a  
 10 compound of the formula 1:



**Formula 1**

wherein **G<sup>1</sup>** is a reactive residue, preferentially a chlorine atom, and the other  
 15 symbols are as defined in general formula I above, with a compound of the formula 2:



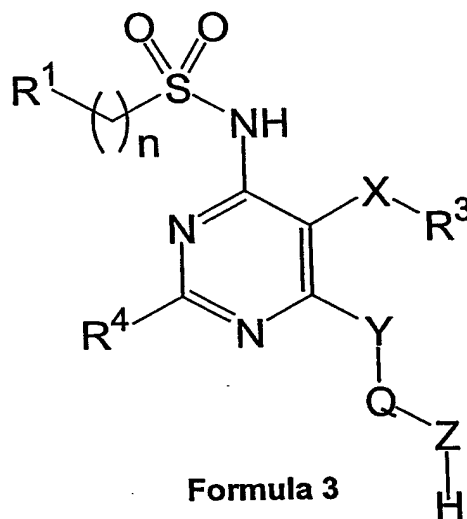
**Formula 2**

20 wherein the symbols are the same as defined in general formula I above, or a salt thereof.

**Possibility B:**

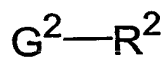
The compounds of general formula I may also be prepared by reacting a compound of formula 3:

5

**Formula 3**

wherein the symbols are the same as defined in general formula I above, or a salt thereof, with a compound of the formula 4

10

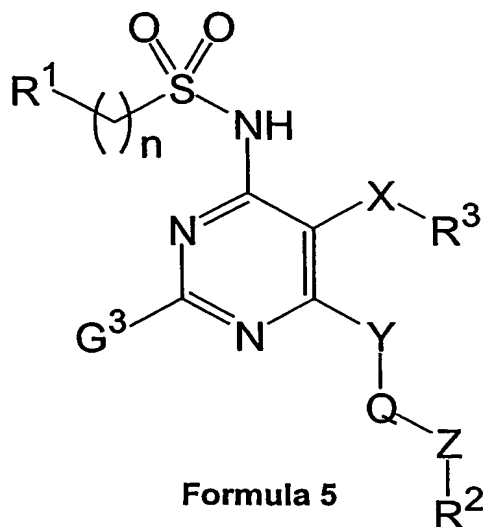
**Formula 4**

wherein  $G^2$  is a reactive residue, such as a halogen atom, and  $R^2$  is as defined in general formula I above.

**Possibility C:**

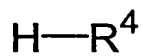
The compounds of general formula I may also be prepared by reacting a compound of the formula 5:

5



Wherein  $G^3$  is a lower alkylsulfonyl group or a phenylsulfonyl group or a halogen atom, and the other symbols are the same as described in general formula I above, or a salt thereof, with a compound of the formula 6:

10

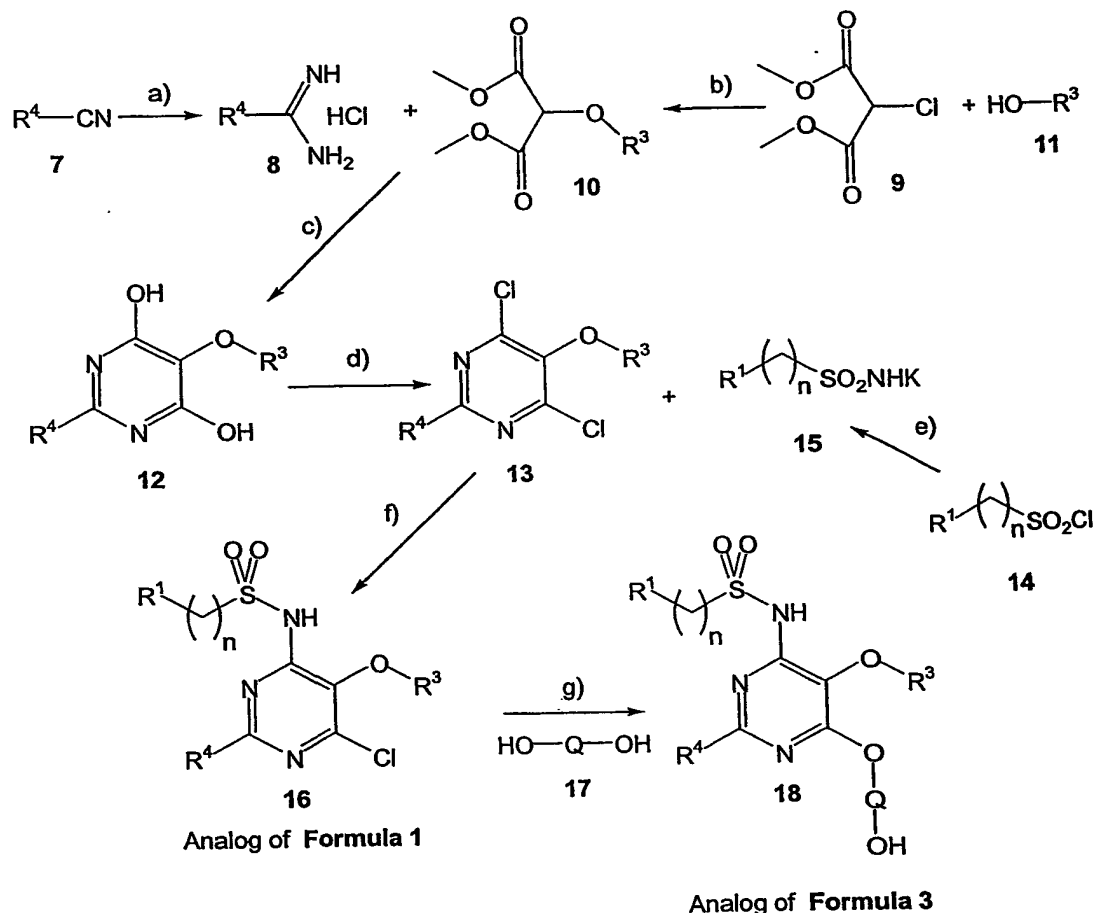
**Formula 6**

wherein  $R^4$  is as defined in general formula I above, or a salt thereof.

15

For possibilities A to C see also [5].

**Scheme 1:** Preparation of the precursors **1** and **3**, with X, Y and Z representing oxygen:



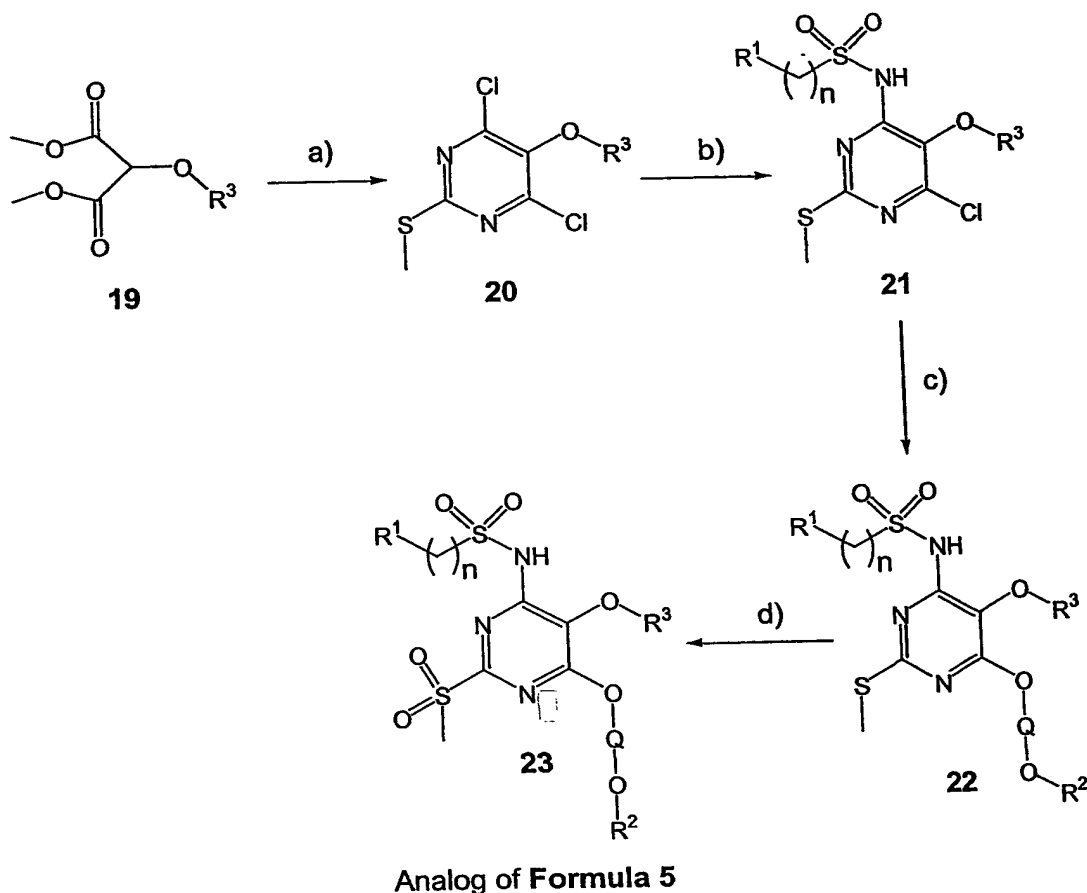
a) NaOMe, MeOH then  $\text{NH}_4\text{Cl}$  or  $\text{LiN}(\text{Si}(\text{CH}_3)_3)_2$  then  $\text{HCl}/i\text{-PrOH}$ ; b)  $\text{K}_2\text{CO}_3$ , acetone; c) NaOMe, MeOH; d)  $\text{POCl}_3$ ; e)  $\text{NH}_3/\text{THF}$  then  $\text{KOtBu}$ , MeOH; f) DMSO; g) NaH, THF, DMF;

- 5 The amidines **8** were synthesized applying standard methodology [1] by reaction of the appropriate nitrile **7** either with sodium methylate in methanol followed by addition of ammonium chloride or by reaction with lithium hexamethyldisilazane followed by addition of hydrochloric acid in *i*-propanol. The 2-substituted malonic esters **10** were prepared according to published procedures [2] by reacting
- 10 dimethylchloromalonate (**9**) with the appropriate alcohol **11** in acetone and potassium carbonate as base. The compounds **10** were dissolved in methanol and sodium methylate was added and stirring was continued for about 30 min followed by the addition of an amidine derivative **8**. Stirring at ambient temperature was continued for another 8 h. After acidic work up the 4,6-

dihydroxypyrimidines **12** could be isolated in yields of 70 to 90% [2]. Compounds **12** or the tautomeric form thereof were transformed into the dichloro derivatives **13** with phosphorus oxychloride in the presence of N,N-dimethylaniline at elevated temperatures (60-120°C) in yields of 40 to 75% [3]. In some cases  
5 better yields were obtained by addition of PCl<sub>5</sub> or benzyl-triethylammoniumchloride. The dichlorides **13** were reacted with an excess of the appropriate sulfonamide potassium salt **15** (prepared according to standard methodology from the sulfochlorides **14** (for the preparation of **14** see e.g. [9], [10]) in DMSO at rt to give the pyrimidines **16** in yields of 70 to 90% either after  
10 recrystallization from EA / diethylether or chromatography through silica gel with EA / heptane. The pyrimidine derivatives **16** are the key intermediates which can be transformed to the desired final products of general formula I either by applying procedures outlined under *Possibility A* or they can be transformed to the derivatives **18** by reaction with a di-hydroxy-compound represented by  
15 formula **17** in the presence of a base such as sodium hydride in a solvent like THF at rt to 90°C and can then be transformed to final compounds according to the general formula I by applying procedures outlined under *Possibility B* above.

20 For further experimental descriptions see [1], [2], [3], [6].  
The synthesis of compounds with X, Y or Z being other groups than oxygen, can be carried out in analogous manners.

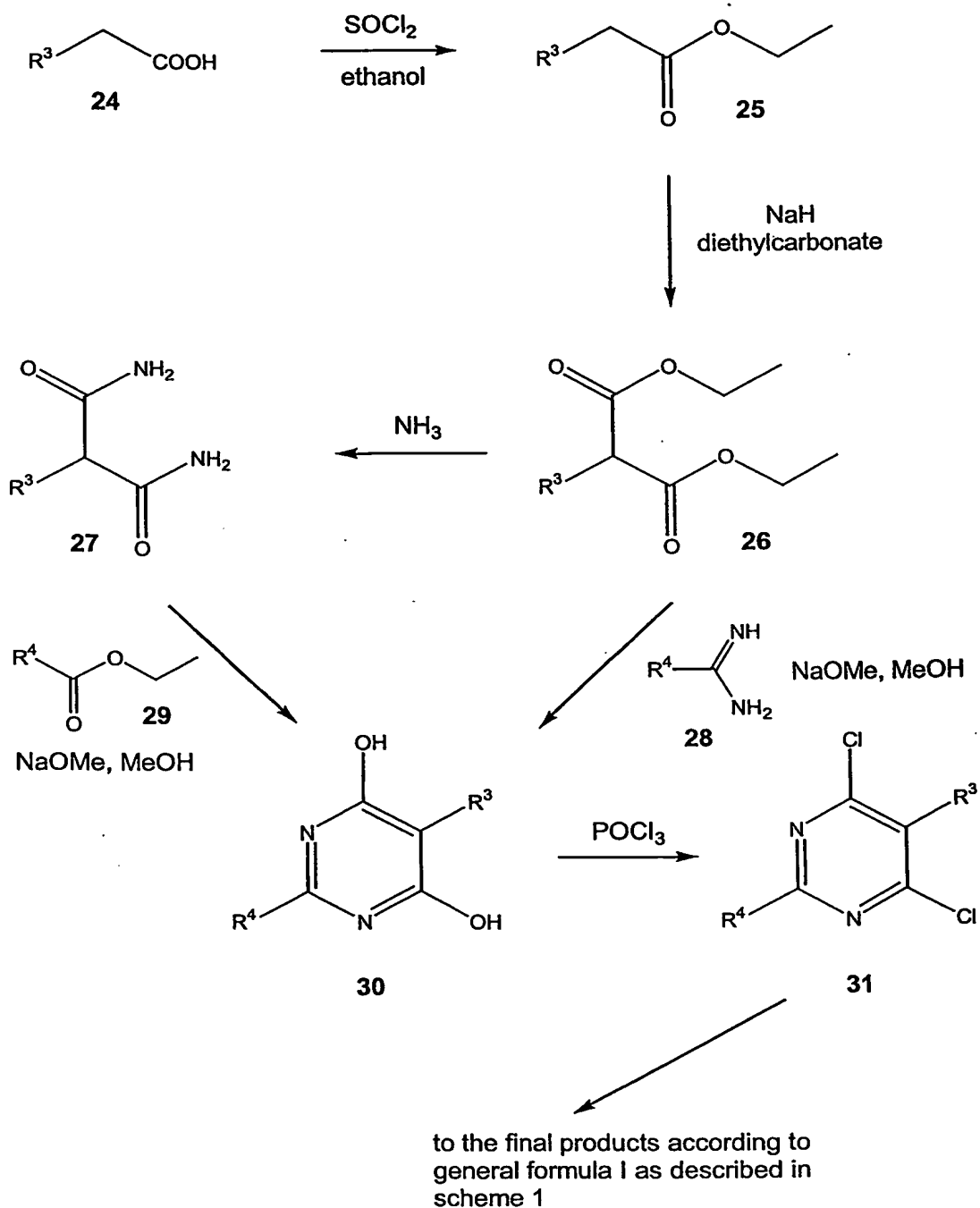
**Scheme 2:** Preparation of the precursor **5**, with X, Y and Z representing oxygen:



- 5 a) i) thiourea, NaOMe, MeOH, rt; ii) CH<sub>3</sub>I, DMSO, rt; iii) POCl<sub>3</sub>, dimethylaniline, 100 – 120°C; b) R<sup>1</sup>-(CH<sub>2</sub>)<sub>n</sub>-SO<sub>2</sub>-NHK, DMSO, rt; c) R<sup>2</sup>-O-Q-OH, NaH, THF/DMF, rt or 60-80°C or HO-Q-OH, NaH, THF/DMF, rt or 60-80°C followed by G<sup>2</sup>-R<sup>2</sup>, NaH, THF, 60-80°C; d) MCPBA, DCM, rt.

- 10 For further experimental descriptions see [1], [2], [3], [5] [6]. For the substitution of the sulfono-group, see especially [5].  
The synthesis of compounds with X, Y or Z being another group than oxygen, can be carried out in analogous procedures.

**Scheme3:** Preparation of the precursors for the synthesis of compounds of general formula I wherein X represents a bond [5]:



In the schemes 1 to 3 the symbols represent the same as defined in general formula I above.

- [1] W. Göhring, J. Schildknecht, M. Federspiel; *Chimia*, **50** (1996), 538 – 543.
- [2] W. Neidhart, V. Breu, D. Bur, K. Burri, M. Clozel, G. Hirth, M. Müller, H. P. Wessel, H. Ramuz; *Chimia*, **50** (1996), 519 – 524 and references cited there.
- 5 [3] W. Neidhart, V. Breu, K. Burri, M. Clozel, G. Hirth, U. Klinkhammer, T. Giller, H. Ramuz; *Bioorg. Med. Chem. Lett.*, **7** (1997), 2223 – 2228. R. A. Nugent, S. T. Schlachter, M. J. Murphy, G. J. Cleek, T. J. Poel, D. G. Whishka, D. R. Graber, Y. Yagi, B. J. Keiser, R. A. Olmsted, L. A. Kopta, S. M. Swaney, S. M. Poppe, J.
- 10 Morris, W. G. Tarpley, R. C. Thomas; *J. Med. Chem.*, **41** (1998), 3793 – 3803.
- [4] J. March; *Advanced Organic Chemistry*, 4<sup>th</sup> Ed., 1994, p. 499 and references cited there.
- 15 [5] EP 0 743 307 A1; EP 0 658 548 B1; EP 0 959 072 A1 (Tanabe Seiyaku)
- [6] EP 0 633 259 B1; EP 0 526 708 A1; WO 96/19459 (F. Hoffmann-LaRoche)
- [7] for the Synthesis of 5-membered heterocycles see: Y. Kohara et al; *J. Med.*
- 20 *Chem.*, **39** (1996), 5228 – 5235 and references cited there.
- [8] EP 0 882 719 A1 (Yamanouchi Pharmaceutical Co., Ltd)
- [9] Z. Zhong, J. A. Bibbs, W. Yuan, C.-H. Wong, *J. Am. Chem. Soc.* **113**, (1991),
- 25 2259-2263
- [10] D. J. Kempf, L. Codavoci, X. C. Wang, W. E. Kohlbrenner, N. E. Wideburg, A. Saldivar, S. Vasavanonda, K. C. Marsh, P. Bryant, H. L. Sham, B. E. Green, D. A. Betebenner, J. Erikson, D. W. Norbeck, *J. Med. Chem.* **36** (1993), 320-330



**Referential Examples (Synthesis of precursors):**

## List of abbreviations:

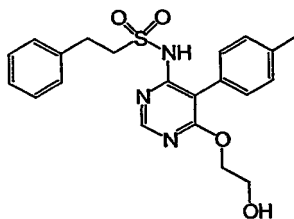
5		
	CyHex	cyclohexane
	DCM	dichloromethane
	DME	1,2-dimethoxyethane
	DMF	dimethylformamide
10	DMSO	dimethylsulfoxide
	EA	ethyl acetate
	Hex	hexane
	HV	high vacuum conditions
	MCPBA	m-chloroperbenzoic acid
15	min	minutes
	rt	room temperature
	THF	tetrahydrofurane
	sat.	saturated
	t <sub>R</sub>	retention time

20

The following referential examples illustrate the invention but do not at all limit the scope thereof.

The following compounds were prepared according to the procedure described above and shown in *Schemes 1 to 3*. All compounds were characterized by <sup>1</sup>H-NMR (300MHz) and occasionally by <sup>13</sup>C-NMR (75MHz) (Varian Oxford, 300MHz; chemical shifts are given in ppm relative to the solvent used; multiplicities: s = singlet, d = doublet, t = triplet; m = multiplett, coupling constants J in Hz), by LC-MS (Waters Micromass; ZMD-platform with ESI-probe with Alliance 2790 HT; 25 Colum: 2x30mm, Gromsil ODS4, 3µm, 120A; Gradient: 0 – 100% acetonitril in water, 6 min, with 0.05% formic acid, flow: 0.45 ml/min; t<sub>R</sub> is given in min.), by 30 TLC (TLC-plates from Merck, Silica gel 60 F<sub>254</sub>) and occasionally by melting point. All temperatures are stated in °C.

## Referential Example 1



5

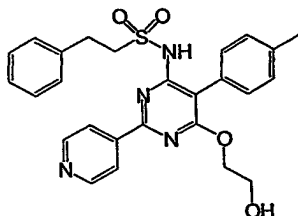
a) Sodium methylate (17 g) was dissolved in methanol (600 ml) at 0°C. 2-p-tolyl-malonic acid diethyl ester (24.5 ml, commercially available from Aldrich), dissolved in 150 ml methanol, was added within 30 min. Stirring was continued for 1 h while slowly warming the mixture to rt. Formamidine hydrochloride (9.9 g, commercially available from Fluka) was added and stirring was continued for 16 h. The solvent was evaporated and 2 M hydrochloric acid (200 ml) was added to the residue followed by slow addition of 10 M sodium hydroxide to adjust the pH to 5. The precipitated product was filtered off and washed subsequently with water and diethylether and dried to give 5-p-tolyl-pyrimidine-4,6-diol (17.7 g). <sup>1</sup>H-NMR (300MHz, d6-DMSO): 8.0(s, 1H); 7.4(d, 2H); 7.1(d, 2H); 2.25(s, 3H).

b) 5-p-tolyl-pyrimidine-4,6-diol (17.2 g) was dissolved in phosphorus oxychloride (250 ml) and N,N-dimethylaniline (25 ml) was added. The mixture was stirred at 70°C for 16 h, then concentrated in vacuo. The residue was poured onto ice-water and extracted with diethylether (3x). The combined organic extracts were washed with 1N hydrochloric acid and with saturated sodium chloride solution, dried over magnesium sulfate, filtered and the filtrate was evaporated. The crude brown material was recrystallized from i-propanol to give 4,6-dichloro-5-p-tolyl-pyrimidine (13.5 g). <sup>1</sup>H-NMR (CDCl<sub>3</sub>): 8.78(s, 1H); 7.35(d, 2H); 7.20(d, 2H); 2.41(s, 3H).

c) 2-Phenylethanesulfonyl chloride was prepared by oxidising phenylethylmercaptan with N-chlorosuccinimide following the procedure given in [9].

- d) A solution of 2-phenylethanesulfonyl chloride (40.94 g) in THF (250 ml) was cooled to  $-20^{\circ}\text{C}$  before it was treated with sat. aq. ammonia (50 ml). The brown suspension was stirred at rt for 16 h. The mixture was neutralised with aq. HCl and the organic solvent was evaporated. The remaining suspension was diluted with water and extracted four times with EA. The organic layers were combined and evaporated to give 2-phenyl-ethanesulfonic acid amide (33.06 g) as orange solid.  $^1\text{H-NMR}$ (300MHz,  $\text{CDCl}_3$ ): 3.15-3.21(m, 2H), 3.38-3.45(m, 2H), 4.59(s br, 2H), 7.21-7.37(m, 5H).
- 10 e) To a solution of 2-phenyl-ethanesulfonic acid amide (33.06 g) in methanol (300 ml) was added K.-tert.butylate (20.03 g). The resulting solution was stirred for 15 min before it was evaporated. The residue was washed with diethyl ether (400 ml) and dried under high vacuum to give 2-phenyl-ethanesulfonic acid amide potassium salt (37.93 g) as an orange powder.
- 15 f) A solution of 2-phenyl-ethanesulfonic acid amide potassium salt (3.0 g), 4,6-dichloro-5-p-tolyl-pyrimidine (2.15 g) and Hunig's base (1.57 ml) in DMSO (50 ml) was stirred at rt for 20 h before it was diluted with water (500 ml) and extracted twice with diethyl ether (250 ml). The aqueous phase was acidified with acetic acid. The resulting suspension was cooled to  $5^{\circ}\text{C}$  and filtered. The solid material was washed with water and diethyl ether, and dried at  $40^{\circ}\text{C}$  under high vacuum to give 2-phenyl-ethanesulfonic acid (6-chloro-5-p-tolyl-pyrimidin-4-yl)-amide (2.08 g) as a grey powder. LC-MS:  $t_R = 5.23$  min,  $[\text{M}+1]^+ = 388.18$ ,  $[\text{M}-1]^- = 386.14$ .
- 20 g) 2-Phenyl-ethanesulfonic acid (6-chloro-5-p-tolyl-pyrimidin-4-yl)-amide (850 mg) was added to a solution of K-tert. butylate (1.1 g) in ethylene glycol (15 ml). The mixture was stirred at  $120^{\circ}\text{C}$  for 27 h before it was diluted with water (100 ml), acidified with 10% aq. citric acid (13 ml). The resulting precipitate was collected, washed with water and diethyl ether and dried to give 2-phenyl-ethanesulfonic acid [6-(2-hydroxy-ethoxy)-5-p-tolyl-pyrimidin-4-yl]-amide (716 mg) as a beige powder. LC-MS:  $t_R = 4.44$  min,  $[\text{M}+1]^+ = 414.18$ ,  $[\text{M}-1]^- = 412.13$ .
- 30

## Referential Example 2



5 a) To a solution of sodium (0.23 g) in methanol (40 ml) was added 4-cyanopyridine (10.62 g) at room temperature. Stirring was continued for 6 h followed by the addition of ammoniumchloride (5.9 g) and stirring was continued for another 10 h. Then diethylether (120 ml) was added and the precipitate was filtered off after 30 min and washed once with diethylether (20 ml). The product was dried under high vacuum. 4-Amidino-pyridine hydrochloride (14.95 g) was  
10 obtained as a white powder.

b) A solution of sodium methylate (6.8 g) in methanol (200 ml) was cooled to 0°C. A solution of diethyl 2-(p-tolyl)-malonate (10.3 g) in methanol (50 ml) was slowly added. Upon completion of the addition the solution was allowed to come  
15 to room temperature and 4-amidino-pyridine hydrochloride (7.57 g) was added. The mixture was stirred at room temperature for 16 h. Eventually, the solvent was removed under reduced pressure and the remaining residue was dissolved in 2 M hydrochloric acid. The solution was extracted with diethyl ether, then adjusted to pH 5 with 10 M sodium hydroxide solution. A precipitate formed. The  
20 precipitate was collected, washed with cold water and dried at 60°C under high vacuum. This gave 4,6-dihydroxy-2-(4-pyridyl)-5-(p-tolyl)-pyrimidine (8.77 g) (or a tautomer) as orange crystals.

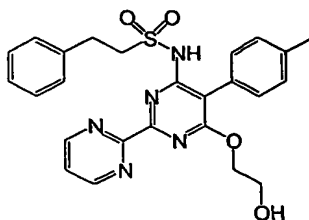
c) To a mixture of 5-(p-tolyl)-4,6-dihydroxy-pyrimidine (8.0 g) and POCl<sub>3</sub> (100 ml),  
25 diethylamine (25 ml) was added at room temperature. The mixture was stirred for 16 h at 60°C. The excess of POCl<sub>3</sub> was distilled off under reduced pressure. The remaining oil was dissolved in DCM (300 ml) and treated with water (300 ml). The aqueous layer was separated and extracted three times with DCM. The combined organic layers were washed with water and brine, dried over MgSO<sub>4</sub> and evaporated. The resulting residue was suspended in isopropanol. The solid  
30

material was collected, washed with isopropanol, and diethyl ether and dried to give 4,6-dichloro-2-(4-pyridyl)-5-(p-tolyl)-pyrimidine (7.2 g) as a white crystalline powder. LC-MS:  $t_R = 5.49$  min,  $[M+1]^+ = 315.89$ .

- 5 d) A solution of 4,6-dichloro-2-(4-pyridyl)-5-(p-tolyl)-pyrimidine (1.5 g), 2-phenyl-ethanesulfonic acid amide potassium salt (1.44 g, Referential Example 1e) and Hunig's base (1 ml) in DMSO (20 ml) was stirred at rt for 24 h before it was diluted with water (150 ml) and extracted twice with diethyl ether. The aqueous layer was acidified with acetic acid. The precipitate was collected, and further  
10 purified by column chromatography on silica gel eluting with hexane:EA 1:1 to give 2-phenyl-ethanesulfonic acid (6-chloro-2-pyridin-4-yl-5-p-tolyl-pyrimidin-4-yl)-amide (480 mg) as a foam. LC-MS:  $t_R = 5.08$  min,  $[M+1]^+ = 465.13$ ,  $[M-1]^- = 462.96$ .
- 15 e) 2-Phenyl-ethanesulfonic acid (6-chloro-2-pyridin-4-yl-5-p-tolyl-pyrimidin-4-yl)-amide (480 mg) was added to a solution of K-tert. butylate (580 mg) in ethylene glycol (5 ml). The mixture was stirred at 110°C for 72 h before it was diluted with water (100 ml), acidified with 10% aq. citric acid (13 ml). The resulting precipitate was collected, washed with water and diethyl ether and dried to give 2-phenyl-  
20 ethanesulfonic acid [6-(2-hydroxy-ethoxy)-2-pyridin-4-yl-5-p-tolyl-pyrimidin-4-yl]-amide as a beige powder. LC-MS:  $t_R = 4.17$  min,  $[M+1]^+ = 491.24$ ,  $[M-1]^- = 489.08$ .

### Referential Example 3

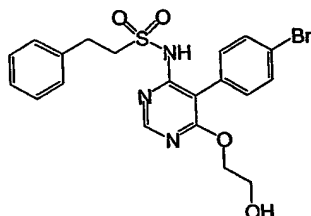
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2-Phenyl-ethanesulfonic acid [6-(2-hydroxy-ethoxy)-5-p-tolyl-[2,2']bipyrimidin-4-yl]-amide was prepared following the procedures given in Referential Example 2

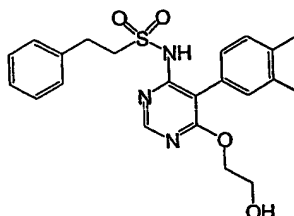
starting from 2-amidino-pyrimidine hydrochloride (e.g. EP 0 526 708 A1) . LC-MS:  $t_R = 4.42$  min,  $[M+1]^+ = 492.30$ ,  $[M-1]^- = 490.27$ .

#### 5 Referential Example 4



2-Phenyl-ethanesulfonic acid [5-(4-bromo-phenyl)-6-(2-hydroxy-ethoxy)-pyrimidin-4-yl]-amide was prepared in analogy to Referential Example 15 using 4-bromophenylacetic acid methyl ester instead of 4-chlorophenylacetic acid methyl ester in step a) and 2-phenyl-ethanesulfonic acid amide potassium salt instead of 2-thiophen-2-yl-ethanesulfonic acid amide potassium salt in step d). LC-MS:  $t_R = 4.60$  min,  $[M+1]^+ = 480.07$ ,  $[M-1]^- = 475.76$ .

#### 15 Referential Example 5

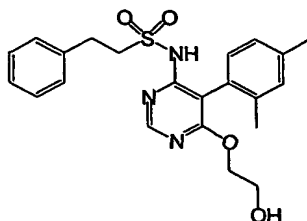


a) (In analogy to a procedure given in *J. Am. Chem. Soc.* **122** (2000), 1360-1370.) To a suspension of  $\text{Pd}(\text{OAc})_2$  (455 mg), 2-(di-tert.-butylphosphino)-biphenyl (1.21 g) and  $\text{K}_3\text{PO}_4$  (39.6 g) in THF (200 ml) dimethylmalonate (12.85 g) and 1-bromo-3,4-dimethyl-benzene 15.0 g) was added under argon. The mixture was refluxed for 16 h, cooled to rt and diluted with EA (300 ml) and filtered. The filtrate was evaporated and the resulting brown oil was purified on silica gel eluting with heptane:EA 4:1 to 1:1 to give 2-(3,4-dimethyl-phenyl)-malonic acid dimethyl ester (16.2 g) as a colourless oil which slowly crystallises.  $^1\text{H-NMR}$

(300MHz, CDCl<sub>3</sub>): 2.25(s, 3H), 2.26(s, 3H), 3.75(s, 6H), 4.59(s, 1H), 7.10-7.20(m, 3H).

- b) 2-Phenyl-ethanesulfonic acid [5-(3,4-dimethyl-phenyl)-6-(2-hydroxy-ethoxy)-pyrimidin-4-yl]-amide was prepared using the above 2-(3,4-dimethyl-phenyl)-malonic acid dimethyl ester in analogy to Referential Example 1. LC-MS:  $t_R$  = 4.61 min,  $[M+1]^+ = 428.19$ ,  $[M-1]^- = 426.07$ .

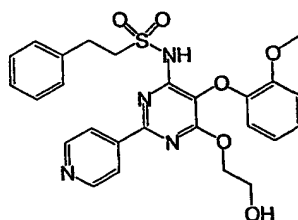
#### 10 Referential Example 6



- a) (In analogy to a procedure given in *J. Am. Chem. Soc.* **122** (2000), 1360-1370.) To a suspension of Pd(OAc)<sub>2</sub> (758 mg), 2-(di-tert.-butylphosphino)-biphenyl (2.02 g) and K<sub>3</sub>PO<sub>4</sub> (65.95 g) in THF (350 ml) dimethylmalonate (21.42 g) and 1-bromo-2,4-dimethyl-benzene (25 g) was added under argon. The mixture was refluxed for 96 h, cooled to rt and diluted with EA (300 ml) and filtered. The filtrate was evaporated and the resulting brown oil was purified on silica gel eluting with heptane:EA 4:1 to 1:1 followed by distillation (bp 95-100°C at 0.064 mbar) to give 2-(2,4-dimethyl-phenyl)-malonic acid dimethyl ester (5.66 g) as a colourless oil. <sup>1</sup>H-NMR(300MHz, CDCl<sub>3</sub>): 2.30(s, 6H), 3.75(s, 6H), 4.87(s, 1H), 6.98-7.05(m, 2H), 7.25-7.28(m, 1H).

- b) 2-Phenyl-ethanesulfonic acid [5-(2,4-dimethyl-phenyl)-6-(2-hydroxy-ethoxy)-pyrimidin-4-yl]-amide was prepared using the above 2-(2,4-dimethyl-phenyl)-malonic acid dimethyl ester in analogy to the procedures given in Referential Example 1. LC-MS:  $t_R$  = 4.54 min,  $[M+1]^+ = 428.23$ ,  $[M-1]^- = 426.07$ .

## Referential Example 7



- 5 a) 2-methoxy-phenol (guaiacol) (48 ml) was slowly added to a stirred suspension of potassium carbonate (70.8 g) in acetone (480 ml) followed by heating to 45°C. Then dimethylchloromalonate (63.2 ml) in acetone (50 ml) was added within 20 min. The reaction mixture was heated to reflux for 16 h. The solvent was evaporated under reduced pressure, the residue taken into water and extracted with DCM. The combined organic layers were dried over sodium sulfate and  
10 evaporated. The oily product was crystallized from methyl-tert.-butyl-ether to give dimethyl-(2-methoxyphenoxy)malonate (86 g).
- b) To a stirred solution of sodium methylate (9.7 g) in methanol (100 ml) a  
15 solution of dimethyl-(2-methoxyphenoxy)malonate (21.7 g) in methanol (50 ml) was added within 15 min and stirring was continued for 30 min followed by the addition of 4-amidino-pyridine hydrochloride (15 g, Referential Example 2) followed by stirring at room temperature for 20 h. The reaction mixture was concentrated in vacuo. The solid residue was stirred with diethyl ether. The  
20 obtained powder was filtered off and dissolved in water (300 ml). Acetic acid was added to pH = 4. The precipitated product was filtered off, washed with water and dried in vacuo at 50°C. 5-(o-methoxyphenoxy)-4,6-dihydroxy-2-(4-pyridyl)-pyrimidine (20.1 g, possibly also present as the tautomeric 5-(2-methoxyphenoxy)-2-(4-pyridyl)-tetrahydropyrimidine-4,6-dion) was obtained as a  
25 white powder.
- c) 5-(2-methoxyphenoxy)-4,6-dihydroxy-2-(4-pyridyl)-pyrimidine (10 g), N-ethyldiisopropylamine (11.2 g), tetraethylammoniumchloride (11 g) and phosphorus pentachloride (13.8 g) were dissolved in phosphorus oxychloride (25  
30 ml) and heated to reflux for 3 h. The mixture was evaporated in vacuo, toluene

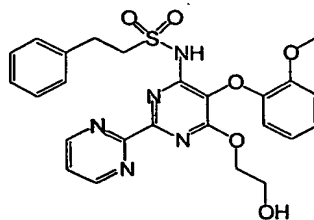


was added and the mixture was again evaporated. The residue was taken into DCM and poured onto ice/water. The layers were separated, the organic layer was washed with water, dried over sodium sulfate and evaporated. After recrystallization from acetone, pure 4,6-dichloro-5-(2-methoxyphenoxy)-2-(4-pyridyl)-pyrimidine (6.52 g) was obtained.

d) A solution of 4,6-dichloro-5-(2-methoxyphenoxy)-2-(4-pyridyl)-pyrimidine (2 g) and 2-phenyl-ethanesulfonic acid amide potassium salt (2.82 g, Referential Example 1e) in DMF (50 ml) was stirred at rt for 16 h. Bulk of the solvent was evaporated before it was diluted with diethyl ether (50 ml). The mixture was acidified with 10% aq. citric acid. The precipitate that formed was collected, washed with diethyl ether (100 ml) and dried to give 2-phenyl-ethanesulfonic acid [6-chloro-5-(2-methoxy-phenoxy)-2-pyridin-4-yl-pyrimidin-4-yl]-amide (2.23 g) as a beige powder. LC-MS:  $t_R = 4.93$  min,  $[M+1]^+ = 497.22$ ,  $[M-1]^- = 494.96$ .

e) To a suspension of NaH (644 mg, 60% in mineral oil) in DME (15 ml) was added ethylene glycol (15 ml). After evolution of gas had ceased, 2-phenyl-ethanesulfonic acid [6-chloro-5-(2-methoxy-phenoxy)-2-pyridin-4-yl-pyrimidin-4-yl]-amide (800 mg) was added and the resulting solution was stirred at 90°C for 16 h. A further portion of NaH (322 mg) was added and stirring was continued at 90°C for 4 d. The mixture was diluted with EA (200 ml) and washed once with 10% aq. citric acid and 3 times with water. The organic phase was evaporated and the residue was suspended in diethyl ether. The solid material was collected, washed with diethyl ether and dried to give 2-phenyl-ethanesulfonic acid [6-(2-hydroxy-ethoxy)-5-(2-methoxy-phenoxy)-2-pyridin-4-yl-pyrimidin-4-yl]-amide (513 mg) as a beige solid. LC-MS:  $t_R = 4.05$  min,  $[M+1]^+ = 523.10$ ,  $[M-1]^- = 521.24$ .

### Referential Example 8

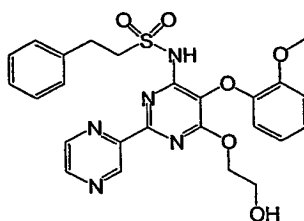


a) 2-Phenylethanesulfonic acid [6-chloro-5-(2-methoxy-phenoxy)-2-(2-pyrimidinyl)-4-pyrimidinyl]-amide was prepared in analogy to Referential Example 7 from 4,6-dichloro-5-(2-methoxyphenoxy)-2-(2-pyrimidinyl)-pyrimidine (prepared as disclosed in [6]) and 2-phenylethane sulfonamide potassium salt (Referential Example 1). LC-MS:  $t_R = 4.85$  min,  $[M+1]^+ = 498.38$ ,  $[M-1]^- = 496.19$ .

b) To a suspension of NaH (803 mg 60% dispersion in mineral oil) in DMF (15 ml) was carefully added ethylene glycol (15 ml). After the evolution of  $H_2$ -gas had stopped 2-phenylethanesulfonic acid [6-chloro-5-(2-methoxy-phenoxy)-2-(2-pyrimidinyl)-4-pyrimidinyl]-amide (1 g) was added. The resulting solution was heated to  $90^\circ C$  and stirred for 16 h. The pale yellow solution was then cooled to rt, diluted with 10% aqueous citric acid solution (100 ml) and extracted three times with EA (50 ml). The combined organic layers were washed once more with 10% aqueous citric acid solution (50 ml) and brine (50 ml) and evaporated. The remaining residue was suspended in water (15 ml). The solid material was filtered off, washed with methanol (50 ml) followed by diethyl ether (50 ml) and dried. This gave 2-phenylethanesulfonic acid [6-(2-hydroxy-ethoxy)-5-(2-methoxy-phenoxy)-2-(2-pyrimidinyl)-4-pyrimidinyl]-amide (766 mg) as a white solid. LC-MS:  $t_R = 4.32$  min,  $[M+1]^+ = 524.47$ ,  $[M-1]^- = 522.29$ .

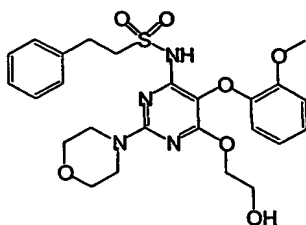
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#### Referential Example 9

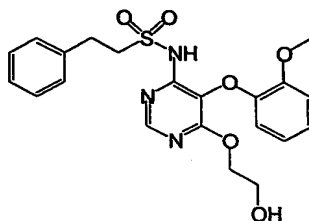


2-Phenyl-ethanesulfonic acid [6-(2-hydroxy-ethoxy)-5-(2-methoxy-phenoxy)-2-pyrazin-2-yl-pyrimidin-4-yl]-amide was prepared following the procedures given in Referential Example 7 starting from 2-amidino-pyrazine hydrochloride. LC-MS:  $t_R = 4.37$  min,  $[M+1]^+ = 524.18$ ,  $[M-1]^- = 522.44$ .

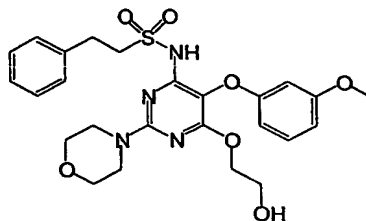
25

**Referential Example 10**

2-Phenyl-ethanesulfonic acid [6-(2-hydroxy-ethoxy)-5-(2-methoxy-phenoxy)-2-morpholin-4-yl-pyrimidin-4-yl]-amide was prepared following the procedures given in Referential Example 7 starting from morpholine-4-carboxamidine hydrobromide. LC-MS:  $t_R = 4.75$  min,  $[M+1]^+ = 531.25$ ,  $[M-1]^- = 529.50$ .

**Referential Example 11**

2-Phenyl-ethanesulfonic acid [6-(2-hydroxy-ethoxy)-5-(2-methoxy-phenoxy)-pyrimidin-4-yl]-amide was prepared following the procedures given in Referential Example 7 starting from formamidine hydrochloride. LC-MS:  $t_R = 4.35$  min,  $[M+1]^+ = 446.15$ ,  $[M-1]^- = 444.11$ .

**Referential Example 12**

a) To a solution of 3-methoxyphenol (115 g) in acetone (1000 ml) was added  $K_2CO_3$  (115 g). The suspension was stirred at 40°C for 15 min. A solution of dimethylchloromalonate (133 ml) in acetone was added over a period of 45 min. The resulting brown suspension was stirred overnight at 70°C. Finally, the solvent was removed under reduced pressure and the residue was taken up in water (1000 ml) and extracted twice with DCM (500 ml). The combined organic

layers were washed with water (500 ml), dried over  $\text{Na}_2\text{SO}_4$ , and evaporated to give crude dimethyl-(3-methoxyphenoxy)malonate (230 g) as an orange oil. The product was not purified any further.

5 b) To a solution of dimethyl-(3-methoxyphenoxy)-malonate (11.19 g) in methanol (100 ml) was added sodium methylate (6.48 g). The yellow solution was stirred for 6 h at rt. Then, morpholine-4-carboxamide hydrobromide (8.40 g) was added and the mixture was stirred at rt for 16 h. The solvent was removed in vacuo and the residue was dissolved in water (150 ml) and extracted twice with  
10 diethyl ether (150 ml). The aqueous phase was acidified with 10% aq. citric acid. The solid that separated was collected, washed with water, evaporated twice from EA, and dried under high vacuum to give 5-(3-methoxy-phenoxy)-2-morpholin-4-yl-pyrimidine-4,6-diol (9.66 g) as a beige powder. LC-MS:  $t_R = 2.88$  min,  $[M+1]^+ = 320.19$ ,  $[M-1]^- = 318.02$ .

15

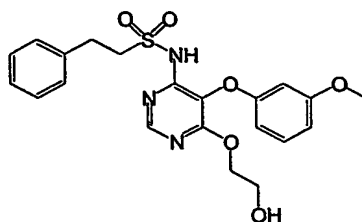
c) 5-(3-Methoxy-phenoxy)-2-morpholin-4-yl-pyrimidine-4,6-diol (9.66 g) was added portionwise to a mixture of  $\text{POCl}_3$  (100 ml) and Hunig's base (50 ml). The black suspension was heated to  $110^\circ\text{C}$  and stirred for 16 h. The mixture was cooled and N,N-dimethylaniline was added before heating was continued for  
20 another 24 h. Bulk of the solvents was evaporated and the remaining oil was poured into water. The dark solution was treated with charcoal before it was extracted twice with EA (300 ml). The organic phase was washed with brine and water, dried over  $\text{MgSO}_4$  and evaporated. The remaining oil was chromatographed on silica gel eluting with heptane:EA. The product was  
25 recrystallised from 2-propanol, the pale yellow crystals were washed with diethyl ether to give 4-[4,6-dichloro-5-(3-methoxy-phenoxy)-pyrimidin-2-yl]-morpholine (7.48 g). LC-MS:  $t_R = 5.56$  min,  $[M+1]^+ = 355.99$ .

d) A solution of 4-[4,6-dichloro-5-(3-methoxy-phenoxy)-pyrimidin-2-yl]-morpholine  
30 (1.0 g) and 2-phenyl-ethanesulfonic acid amide potassium salt (1.57 g, Referential Example 1e) in DMSO (15 ml) was stirred at  $60^\circ\text{C}$  for 24 h. The solution was diluted with water (75 ml) and extracted twice with diethyl ether (75 ml) before it was acidified with 10% aq. citric acid. The mixture was extracted

twice with EA (150 ml). The organic phase was washed with water (50 ml). The product precipitated upon evaporation of the solvent. The solid was collected, washed with diethyl ether and dried to give 2-phenyl-ethanesulfonic acid [6-chloro-5-(3-methoxy-phenoxy)-2-morpholin-4-yl-pyrimidin-4-yl]-amide (1.28 g) as  
5 off-white powder. LC-MS:  $t_R = 5.34$  min,  $[M+1]^+ = 505.12$ ,  $[M-1]^- = 502.97$ .

e) A suspension of 2-phenyl-ethanesulfonic acid [6-chloro-5-(3-methoxy-phenoxy)-2-morpholin-4-yl-pyrimidin-4-yl]-amide (1.27 g) in ethylene glycol (12 ml) was treated with K-tert.-butylate (2.82 g). The resulting solution was stirred at  
10  $100^\circ\text{C}$  for 12 d. The solution was cooled to rt, diluted with 10% aq. citric acid (150 ml) and extracted twice with EA (150 ml). The organic phase was washed with water (50 ml) and evaporated. The crude product was purified by chromatography on silica gel eluting with heptane:EA 1:1 to 1:2 to furnish 2-phenyl-ethanesulfonic acid [6-(2-hydroxy-ethoxy)-5-(3-methoxy-phenoxy)-2-morpholin-4-yl-pyrimidin-4-yl]-amide (1.1 g) as a white solid. LC-MS:  $t_R = 4.66$   
15 min,  $[M+1]^+ = 531.20$ ,  $[M-1]^- = 529.14$ .

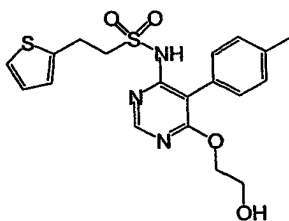
### Referential Example 13



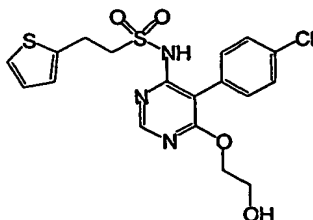
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2-Phenyl-ethanesulfonic acid [6-(2-hydroxy-ethoxy)-5-(3-methoxy-phenoxy)-pyrimidin-4-yl]-amide was prepared following the procedures given in Referential Example 12 starting from formamidine hydrochloride.  $^1\text{H-NMR}$  (300MHz,  $\text{CDCl}_3$ ): 3.14-3.21(m, 2H), 3.70-3.74(m, 2H), 3.79(s, 3H), 3.94-4.01(m, 2H), 4.40-4.46(m, 2H), 6.41(dd,  $J=2.4, 8.4$ , 1H), 6.47(t,  $J=2.4$ , 1H), 6.66(dd,  $J=1.8, 8.4$ , 1H), 7.16-7.32(m, 6H), 8.37(s, 1H); LC-MS:  $t_R = 4.33$  min,  $[M+1]^+ = 446.27$ ,  $[M-1]^- = 444.05$ .  
25

## Referential Example 14



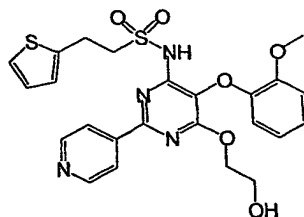
- a) 2-Thiophen-2-yl-ethanesulfonyl chloride was obtained starting from commercial 2-(2-bromo-ethyl)-thiophene following the procedures given in the literature (J. Am. Chem. Soc. 103 (1981), 1525-1533).
- b) A solution of crude 2-thiophen-2-yl-ethanesulfonyl chloride (25 g) in THF (400 ml) was treated with sat. aq. ammonia (60 ml) at 0°C. The mixture was stirred at rt for 16 h before it was neutralised with 25% aq. HCl (60 ml). Bulk of the THF was evaporated. The aq. solution was extracted twice with EA. The organic phase was washed with water and evaporated. The remaining oil was purified by chromatography on silica gel eluting with heptane:EA 1:1. The product was further purified by recrystallisation from diethyl ether/pentane to give 2-thiophen-2-yl-ethanesulfonic acid amide (8.46 g) as off-white crystals. <sup>1</sup>H-NMR (300MHz, CDCl<sub>3</sub>): 3.36-3.50(m, 4H), 4.54(s br, 2H), 6.89-6.92(m, 1H), 6.95(dd, J=3.5, 5.1, 1H), 7.20(dd, J=1.1, 5.0, 1H).
- c) A solution of 2-thiophen-2-yl-ethanesulfonic acid amide (3.77 g) in methanol (200 ml) was treated with K-tert.-butylate (2.21 g), stirred at rt for 15 min, evaporated and dried under high vacuum to give 2-thiophen-2-yl-ethanesulfonic acid amide potassium salt (4.5 g) as a beige powder.
- d) 2-Thiophen-2-yl-ethanesulfonic acid [6-(2-hydroxy-ethoxy)-5-p-tolyl-pyrimidin-4-yl]-amide was prepared using the above 2-thiophen-2-yl-ethanesulfonic acid amide potassium salt following the procedures given in Referential Example 1. <sup>1</sup>H-NMR(300MHz, CDCl<sub>3</sub>): 2.42(s, 3H), 3.33-3.41(m, 2H), 3.81-3.86(m, 2H), 3.99-4.07(m, 2H), 4.46-4.51(m, 2H), 6.82-6.86(m, 1H), 6.92(dd, J=3.5, 5.1, 1H), 7.13-7.18(m, 3H), 7.28-7.32(m 2H), 8.51(s, 1H); LC-MS: t<sub>R</sub> = 4.18 min, [M+1]<sup>+</sup> = 420.24, [M-1]<sup>-</sup> = 418.20.

**Referential Example 15**

- 5 a) At 35°C a solution of 4-chlorophenylacetic acid methyl ester (52 g) in THF (170 ml) was carefully added over a period of 70 min to a suspension of NaH (15.6 g) in dry THF (550 ml) . Stirring was continued for 40 min without heating and the temperature dropped to 29°C. The evolution of gas had stopped before dimethylcarbonate (94.8 ml) was added dropwise while the temperature of the
- 10 mixture was maintained at 25-28°C. After the evolution of gas had ceased, the mixture was diluted with THF (200 ml) and stirring was continued for 72 h at rt. The mixture was carefully acidified with aq. HCl before bulk of the THF was removed in vacuo. The residue was dissolved in diethyl ether (1200 ml), washed three times with 1 N aq. HCl and once with brine, dried over MgSO<sub>4</sub> and
- 15 evaporated. The residue formed was collected, washed with diethyl ether and dried to give 2-(4-chloro-phenyl)-malonic acid dimethyl ester (42 g) as white crystals.
- b) A solution of 2-(4-chlorophenyl)-malonic acid dimethyl ester (18.90 g) in
- 20 methanol (200 ml) was added dropwise at 0°C to a solution sodium methylate (14.60 g) in methanol (150 ml). The mixture was stirred for 1 h at 0°C before formamidine hydrochloride (7.66 g) was added. The suspension was stirred at rt for 20 h. The solvent was removed and the residue was suspended in 2 N aq. HCl (200 ml). The pH of the suspension was carefully adjusted to 4-5 by adding
- 25 10 M NaOH (20 ml), stirring was continued for 30 min. The white precipitate was collected, washed with water and diethyl ether and dried to give 5-(4-chlorophenyl)-pyrimidine-4,6-diol (16.44 g) as a white powder. LC-MS:  $t_R$  = 2.75 min,  $[M+H]^+$  = 222.96,  $[M-H]^-$  = 220.92.

- c) To a suspension of 5-(4-chlorophenyl)-pyrimidine-4,6-diol (16.44 g) in  $\text{POCl}_3$  (165 ml) was carefully added N, N-dimethylaniline (16.5 ml). The mixture was refluxed for 1.5 h. The dark green solution was evaporated and the residue was poured onto ice/water. The suspension was diluted with 2 N HCl (200 ml) and water (800 ml) and stirred at  $2^\circ\text{C}$  for 1 h. The precipitate was collected, washed with water and dried to give 4,6-dichloro-5-(4-chlorophenyl)-pyrimidine (18.66 g) as a slightly green powder.
- d) A solution of 4,6-dichloro-5-(4-chlorophenyl)-pyrimidine (848 mg), 2-thiophen-2-yl-ethanesulfonic acid amide potassium salt (1.5 g, Referential Example 14) and Hunig's base (1 ml) in DMSO (20 ml) was stirred at rt for 24 h before it was diluted with water (200 ml) and extracted twice with diethyl ether. The aqueous layer was acidified with acetic acid. The precipitate was collected, washed with water and diethyl ether and dried to give 2-thiophen-2-yl-ethanesulfonic acid [6-chloro-5-(4-chloro-phenyl)-pyrimidin-4-yl]-amide (930 mg) as a beige powder. LC-MS:  $t_R = 5.01$  min,  $[\text{M}+1]^+ = 413.49$ ,  $[\text{M}-1]^- = 411.93$ .
- e) 2-Thiophen-2-yl-ethanesulfonic acid [6-chloro-5-(4-chloro-phenyl)-pyrimidin-4-yl]-amide (930 mg) was added to a solution of K-tert. butylate (1.16 g) in ethylene glycol (10 ml). The mixture was stirred at  $110^\circ\text{C}$  for 12 h before it was diluted with water (150 ml), acidified with 10% aq. citric acid (13 ml). The resulting precipitate was collected, washed with water and diethyl ether and dried to give (820 mg) 2-thiophen-2-yl-ethanesulfonic acid [5-(4-chloro-phenyl)-6-(2-hydroxy-ethoxy)-pyrimidin-4-yl]-amide as a beige powder. LC-MS:  $t_R = 4.43$  min,  $[\text{M}+1]^+ = 440.01$ ,  $[\text{M}-1]^- = 437.99$ .

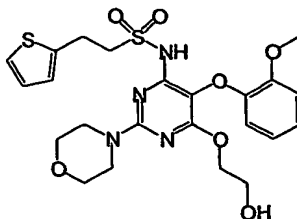
### Referential Example 16





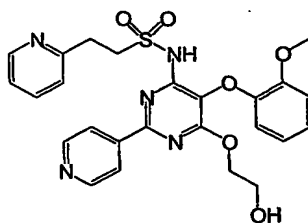
2-Thiophen-2-yl-ethanesulfonic acid [6-(2-hydroxy-ethoxy)-5-(2-methoxy-phenoxy)-2-pyridin-4-yl-pyrimidin-4-yl]-amide was prepared following the procedures given in Referential Example 7 using 2-thiophen-2-yl-ethanesulfonic acid amide potassium salt (Referential Example 14).; LC-MS:  $t_R = 4.00$  min,  $[M+1]^+ = 529.29$ ,  $[M-1]^- = 526.97$ .

#### Referential Example 17



2-Thiophen-2-yl-ethanesulfonic acid [6-(2-hydroxy-ethoxy)-5-(2-methoxy-phenoxy)-2-morpholin-4-yl-pyrimidin-4-yl]-amide was prepared following the procedures given in Referential Example 10 using 2-thiophen-2-yl-ethanesulfonic acid amide potassium salt (Referential Example 14).; LC-MS:  $t_R = 4.62$  min,  $[M+1]^+ = 537.21$ ,  $[M-1]^- = 534.96$ .

#### Referential Example 18



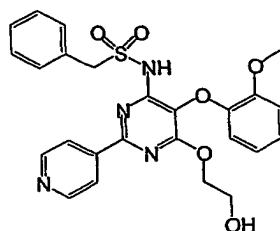
a) 2-Pyridin-2-yl-ethanesulfonyl chloride hydrochloride was prepared starting from commercial 2-pyridine-2-ethane sulfonic acid following the procedure given in *J. Med. Chem.* 36 (1993), 320-330.

b) 2-Pyridin-2-yl-ethanesulfonic acid amide potassium salt was prepared using the above 2-pyridin-2-yl-ethanesulfonyl chloride hydrochloride following the procedures given in Referential Example 14 b and 14c.

c) A solution of 4,6-dichloro-5-(2-methoxyphenoxy)-2-(4-pyridyl)-pyrimidine (1.75 g, Referential Example 7) and 2-pyridin-2-yl-ethanesulfonic acid amide potassium salt (1.13 g) in DMSO (30 ml) was stirred at rt for 24 h. Triethylamine (657 mg) was added and stirring was continued for another 96 h before the mixture was diluted with ethyl acetate (150 ml) and washed 4% aq. citric acid and water. The aqueous phase was extracted three more times with EA. The organic phase was dried over  $\text{MgSO}_4$  and evaporated. The crude product was purified by column chromatography on silica gel eluting with EA containing 0-10% methanol to give 2-pyridin-2-yl-ethanesulfonic acid [6-chloro-5-(2-methoxyphenoxy)-2-pyridin-4-yl-pyrimidin-4-yl]-amide (950 mg) as a brown powder. LC-MS:  $t_R = 3.63$  min,  $[\text{M}+1]^+ = 498.31$ ,  $[\text{M}-1]^- = 496.10$ .

d) To a suspension of NaH (701 mg, 60% in mineral oil) in DMF (15 ml) was added ethylene glycol (15 ml). After the evolution of gas had ceased, 2-pyridin-2-yl-ethanesulfonic acid [6-chloro-5-(2-methoxyphenoxy)-2-pyridin-4-yl-pyrimidin-4-yl]-amide (800 mg) was added and the resulting solution was stirred at  $90^\circ\text{C}$  for 40 h. The solution was neutralised with 2 N aq. HCl (7 ml) before it was evaporated. The brown residue was purified by chromatography on prep. tlc-plates with EA:methanol:sat. aq. ammonia 10:2:1. The product was further purified by crystallisation from methanol:diethyl ether:pentane to give 2-pyridin-2-yl-ethanesulfonic acid [6-(2-hydroxy-ethoxy)-5-(2-methoxyphenoxy)-2-pyridin-4-yl-pyrimidin-4-yl]-amide (280 mg) as beige crystals.  $^1\text{H-NMR}$ (300MHz,  $\text{CDCl}_3$ ): 3.36-3.43(m, 2H), 3.86-3.90(m, 2H), 3.94(s, 3H), 4.23-4.30(m, 2H), 4.56-4.62(m, 2H), 6.91(dt,  $J_d=1.5$ ,  $J_t=7.7$ , 1H, 7.00(dd,  $J=1.7$ , 8.2, 1H, 7.08-7.20(m, 4H), 7.57(dt,  $J_d=1.8$ ,  $J_t=7.9$ , 1H), 8.15(dd,  $J=1.7$ , 4.6, 2H), 8.43(d,  $J=4.4$ , 1H), 8.72(dd,  $J=1.7$ , 4.6, 2H); LC-MS:  $t_R = 3.23$  min,  $[\text{M}+1]^+ = 524.48$ ,  $[\text{M}-1]^- = 522.25$ .

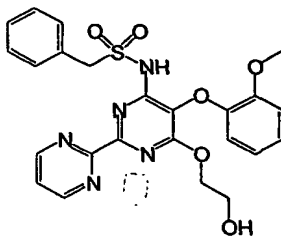
### Referential Example 19



a) Phenyl-methanesulfonamide potassium salt was prepared in analogy to the procedures given in Referential Example 1d and 1e using commercial phenyl-methanesulfonyl chloride.  $^1\text{H-NMR}$ (300MHz, DMSO): 3.73(s, 2H), 7.13-7.30(m, 5H).

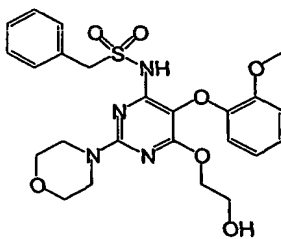
b) N-[6-(2-Hydroxy-ethoxy)-5-(2-methoxy-phenoxy)-2-pyridin-4-yl-pyrimidin-4-yl]-C-phenyl-methanesulfonamide was prepared in analogy to Referential Example 7 using the above phenyl-methanesulfonamide potassium salt. LC-MS:  $t_R = 3.99$  min,  $[\text{M}+1]^+ = 509.32$ ,  $[\text{M}-1]^- = 507.31$ .

#### Referential Example 20



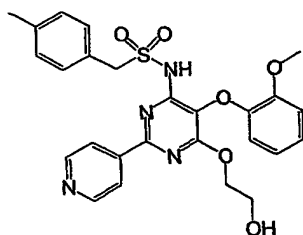
N-[6-(2-Hydroxy-ethoxy)-5-(2-methoxy-phenoxy)-[2,2']bipyrimidin-4-yl]-C-phenyl-methanesulfonamide was prepared in analogy to Referential Example 8 using phenyl-methanesulfonamide potassium salt (Referential Example 19). LC-MS:  $t_R = 4.15$  min,  $[\text{M}+1]^+ = 510.34$ ,  $[\text{M}-1]^- = 508.54$ .

#### Referential Example 21



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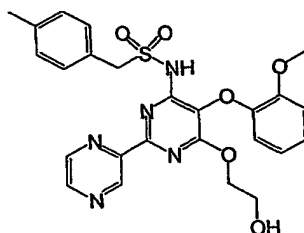
N-[6-(2-Hydroxy-ethoxy)-5-(2-methoxy-phenoxy)-2-morpholin-4-yl-pyrimidin-4-yl]-C-phenyl-methanesulfonamide was prepared in analogy to Referential Example 10 using phenyl-methanesulfonamide potassium salt (Referential Example 19). LC-MS:  $t_R = 4.54$  min,  $[\text{M}+1]^+ = 517.32$ ,  $[\text{M}-1]^- = 515.07$ .

**Referential Example 22**

a) p-Tolyl-methanesulfonyl chloride was prepared by oxidising commercially  
5 available p-tolyl-methanethiol with N-chlorosuccinimide in analogy to the  
procedure disclosed in [9].

b) p-Tolyl-methanesulfonamide potassium salt was prepared in analogy to the  
procedures given in Referential Example 1d and 1e. <sup>1</sup>H-NMR(300MHz, CDCl<sub>3</sub>)  
10 (sulfonamide): 2.36(s, 3H), 4.27(s, 2H), 4.63(s br, 2H), 7.20(d, J=7.9, 2H),  
7.30(d, J=8.1, 2H).

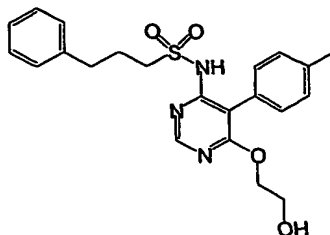
c) N-[6-(2-Hydroxy-ethoxy)-5-(2-methoxy-phenoxy)-2-pyridin-4-yl-pyrimidin-4-yl]-  
C-p-tolyl-methanesulfonamide was prepared using the above p-tolyl-  
15 methanesulfonamide potassium salt in analogy to the procedures given in  
Referential Example 7. LC-MS: t<sub>R</sub> = 4.18 min, [M+1]<sup>+</sup> = 523.22, [M-1]<sup>-</sup> = 521.19.

**Referential Example 23**

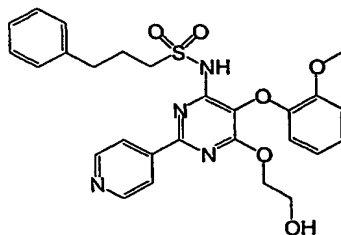
20

N-[6-(2-Hydroxy-ethoxy)-5-(2-methoxy-phenoxy)-2-pyrazin-2-yl-pyrimidin-4-yl]-C-  
p-tolyl-methanesulfonamide was prepared using the above p-tolyl-  
methanesulfonamide potassium salt in analogy to the procedures given in  
Referential Example 9. LC-MS: t<sub>R</sub> = 4.46 min, [M+1]<sup>+</sup> = 524.20, [M-1]<sup>-</sup> = 521.93.

25

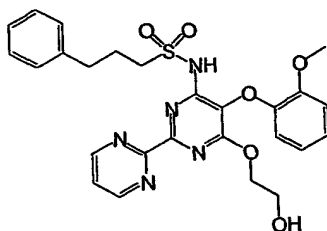
**Referential Example 24**

- a) 3-Phenyl-propane-1-sulfonyl chloride was prepared by oxidising commercially available 3-phenyl-propane-1-thiol with N-chlorosuccinimide in analogy to the procedure disclosed in [9].
- b) 3-Phenyl-propane-1-sulfonic acid amide potassium salt was prepared in analogy to the procedures given in Referential Example 1d and 1e. <sup>1</sup>H-NMR (300MHz, CDCl<sub>3</sub>) (sulfonamide): 2.11-2.23(m, 2H), 2.76(t, J=7.5, 2H), 3.05-3.13(m, 2H), 4.85(s br, 2H), 7.14-7.40(m, 5H).
- c) 3-Phenyl-propane-1-sulfonic acid [6-(2-hydroxy-ethoxy)-5-p-tolyl-pyrimidin-4-yl]-amide was prepared in analogy to the procedures given in Referential Example 1 using the above 3-phenyl-propane-1-sulfonic acid amide potassium salt. LC-MS: t<sub>R</sub> = 4.66 min, [M+1]<sup>+</sup> = 428.24, [M-1]<sup>-</sup> = 426.21.

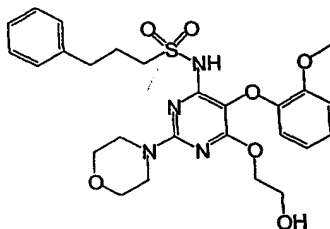
**Referential Example 25**

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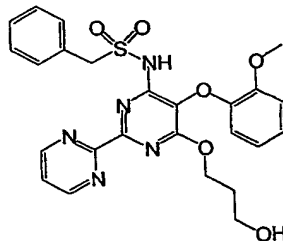
- 3-Phenyl-propane-1-sulfonic acid [6-(2-hydroxy-ethoxy)-5-(2-methoxy-phenoxy)-2-pyridin-4-yl-pyrimidin-4-yl]-amide was prepared in analogy to the procedures given in Referential Example 7 using 3-phenyl-propane-1-sulfonic acid amide potassium salt of Referential Example 24. LC-MS: t<sub>R</sub> = 4.14 min, [M+1]<sup>+</sup> = 537.45, [M-1]<sup>-</sup> = 535.41.
- 25

**Referential Example 26**

3-Phenyl-propane-1-sulfonic acid [6-(2-hydroxy-ethoxy)-5-(2-methoxy-phenoxy)-  
5 [2,2']bipyrimidin-4-yl]-amide was prepared in analogy to the procedures given in  
Referential Example 8 using 3-phenyl-propane-1-sulfonic acid amide potassium  
salt of Referential Example 24. LC-MS:  $t_R = 4.37$  min,  $[M+1]^+ = 538.38$ ,  $[M-1]^- =$   
536.27.

**10 Referential Example 27**

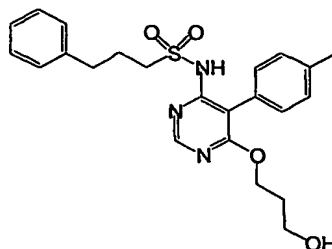
3-Phenyl-propane-1-sulfonic acid [6-(2-hydroxy-ethoxy)-5-(2-methoxy-phenoxy)-  
2-morpholin-4-yl-pyrimidin-4-yl]-amide was prepared in analogy to the  
15 procedures given in Referential Example 10 using 3-phenyl-propane-1-sulfonic  
acid amide potassium salt of Referential Example 24. LC-MS:  $t_R = 4.80$  min,  
 $[M+1]^+ = 545.40$ ,  $[M-1]^- = 543.52$ .

**Referential Example 28**

N-[6-(3-Hydroxy-propoxy)-5-(2-methoxy-phenoxy)-[2,2']bipyrimidinyl-4-yl]-C-phenyl-methanesulfonamide was prepared in analogy to the procedures given in Referential Example 20 using propane-1,3-diol instead of ethylene glycol. LC-MS:  $t_R = 4.13$  min,  $[M+1]^+ = 524.34$ ,  $[M-1]^- = 522.19$ .

5

### Referential Example 29



3-Phenyl-propane-1-sulfonic acid [6-(3-hydroxy-propoxy)-5-p-tolyl-pyrimidin-4-yl]-amide was prepared in analogy to the procedures given in Referential Example 24 using propane-1,3-diol instead of ethylene glycol. LC-MS:  $t_R = 4.72$  min,  $[M+1]^+ = 442.28$ ,  $[M-1]^- = 440.22$ .

### Examples

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The following examples illustrate the invention but do not at all limit the scope thereof.

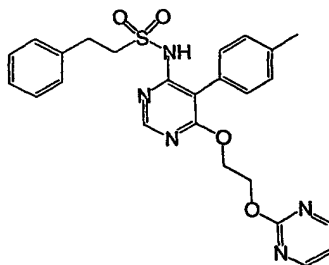
The following reagents have been synthesised in order to prepare the examples listed below according to or in analogy to literature procedures: 5-bromo-2-chloro-pyrimidine (*Aust. J. Chem.* **17** (1964), 794-802; *J. Org. Chem.* **25** (1960), 1916-1919); 2,5-dichloro-pyrimidine (in analogy to 5-bromo-2-chloro-pyrimidine using chlorine instead of bromine); 2-chloro-5-methyl-pyrimidine (*J. Med. Chem.* **6** (1963), 697-701; *Aust. J. Chem.* **30** (1977), 2515-2525); 2-methanesulfonyl-5-methoxy-pyrimidine (*J. Chem. Soc. Perkin Trans. 1*, **1999**, 3265-3268; *J. Org. Chem.* **27** (1962), 3614-3617); 2-chloro-5-methylsulfanyl-pyrimidine (*Il Farmaco* **43** (1988), 277-292; French Patent 1 549 494 (1968)); 2-methanesulfonyl-5-trifluoromethyl-pyrimidine (*Tetrahedron Lett.* **37** (1996), 1829-1832); 2-

methanesulfonyl-4,6-dimethoxy-pyrimidine was prepared starting from 4,6-dichloro-2-methylsulfanyl-pyrimidine using standard methodology.

All other reagents constitute commercially available ingredients.

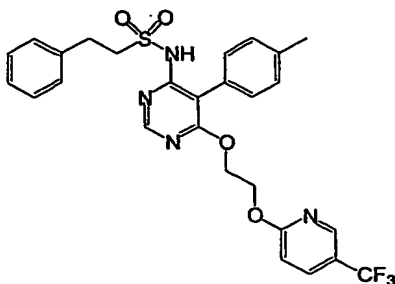
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### Example 1



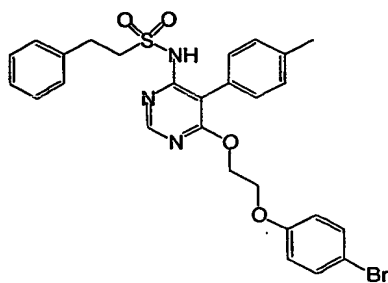
- 10 To sodium hydride (50 mg, 60% in mineral oil) was added THF (25 ml) followed by 2-phenyl-ethanesulfonic acid [6-(2-hydroxy-ethoxy)-5-p-tolyl-pyrimidin-4-yl]-amide (150 mg, Referential Example 1). The mixture was stirred for 1 h at rt before 2-chloro-pyrimidine (86 mg) was added. Stirring was continued for 17 h at 80°C. The solvent was evaporated and diethylether (20 ml) was added to the
- 15 residue. The precipitate was filtered off and washed with diethylether, dissolved in water (20 ml) and acidified with citric acid and extracted twice with EA (50 ml). The organic phase was washed with water, dried over Na<sub>2</sub>SO<sub>4</sub> and evaporated. The residue is suspended in 2-propanol (15 ml) and stirred at 70°C for 10 min and cooled to 0°C before the solid material is collected, washed with 2-propanol
- 20 (2 ml) and dried under high vacuum to furnish 2-phenyl-ethanesulfonic acid {6-[2-(pyrimidin-2-yloxy)-ethoxy]-5-p-tolyl-pyrimidin-4-yl}-amide (133 mg) as a white powder. LC-MS:  $t_R = 4.97$  min,  $[M+1]^+ = 492.34$ ,  $[M-1]^- = 490.28$ .



**Example 2**

- 5 To sodium hydride (27 mg, 60% in mineral oil) was added THF (15 ml) followed by 2-phenyl-ethanesulfonic acid [6-(2-hydroxy-ethoxy)-5-p-tolyl-pyrimidin-4-yl]-amide (80 mg, Referential Example 1). The mixture was stirred for 1 h at rt before 2-chloro-5-trifluoromethyl-pyridine (86 mg) was added. Stirring was continued for 17 h at 80°C. The solvent was evaporated, the residue was
- 10 dissolved in water (20 ml), acidified with citric acid. The suspension was treated with hexane (15 ml). The solid material was collected, washed with hexane:EA 1:1 (20 ml) and dried to yield 2-phenyl-ethanesulfonic acid {5-p-tolyl-6-[2-(5-trifluoromethyl-pyridin-2-yloxy)-ethoxy]-pyrimidin-4-yl}-amide (96 mg) as beige powder. LC-MS:  $t_R = 5.86$  min,  $[M+1]^+ = 559.20$ ,  $[M-1]^- = 557.37$ .

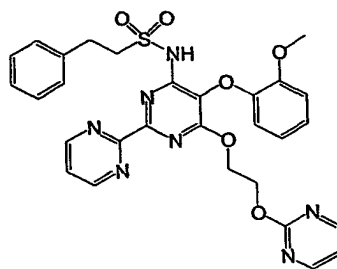
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**Example 3**

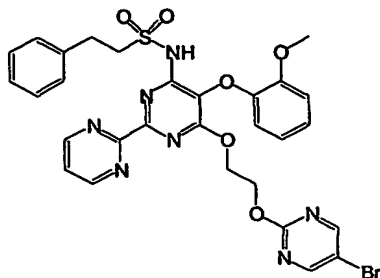
- 20 To a suspension of NaH (17 mg, 60% in mineral oil) in DME (5 ml) was added 2-(4-bromo-phenoxy)-ethanol (111 mg). The mixture was stirred at 50°C for 1h before 2-phenyl-ethanesulfonic acid (6-chloro-5-p-tolyl-pyrimidin-4-yl)-amide (100 mg, Referential Example 1f) and K-tert.-butylate (25 mg) was added. The mixture was stirred at 70°C for 16 h. A further portion of K-tert.-butylate (50 mg) was

added and stirring was continued at 70°C for 12 h and at rt for 72 h. The solvent was evaporated. The residue was treated with water (40 ml), acidified with 10% aq. citric acid and extracted twice with EA (50 ml). The organic phase was washed with water and evaporated. The crude product was crystallised from 2-propanol to give 2-phenyl-ethanesulfonic acid {6-[2-(4-bromo-phenoxy)-ethoxy]-5-p-tolyl-pyrimidin-4-yl}-amide (127 mg) as a beige powder. LC-MS:  $t_R$  = 6.14 min,  $[M+1]^+ = 568.38$ ,  $[M-1]^- = 570.15$ .

#### Example 4

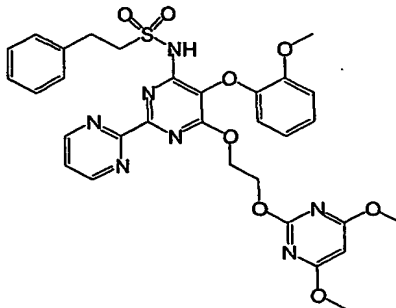


To a suspension of NaH (29 mg 60% dispersion in mineral oil) in a mixture of DMF and THF (2.5 ml each) was added 2-phenylethanesulfonic acid [6-(2-hydroxy-ethoxy)-5-(2-methoxy-phenoxy)-2-(2-pyrimidinyl)-4-pyrimidinyl]-amide (150 mg). After gas-evolution had ceased, 2-chloropyrimidine (82 mg) was added. The resulting pale yellow suspension was stirred at 70°C for 4 h. It was cooled to rt, diluted with EA (75 ml) and washed with 10% aqueous citric acid solution (50 ml) followed by water (50 ml). The organic layer was evaporated and the remaining residue was purified on prep. tlc plates (silica gel, 0.5 mm layer thickness) developing with EA:methanol:sat. aqueous ammonia 8:2:1. This furnished 2-phenylethanesulfonic acid {5-(2-methoxy-phenoxy)-6-[2-(pyrimidin-2-yloxy)-ethoxy]-2-(2-pyrimidinyl)-4-pyrimidinyl}-amide as a pale yellow powder. LC-MS:  $t_R$  = 4.60 min,  $[M+1]^+ = 602.69$ ,  $[M-1]^- = 600.43$ .

**Example 5**

- 5 To a suspension of NaH (29 mg 60% dispersion in mineral oil) in a mixture of DMF and THF (2.5 ml each) was added 2-phenylethanesulfonic acid [6-(2-hydroxy-ethoxy)-5-(2-methoxy-phenoxy)-2-(2-pyrimidinyl)-4-pyrimidinyl]-amide (150 mg). After gas-evolution had ceased, 2-chloro-5-bromopyrimidine (138 mg) was added. The resulting orange suspension was stirred at 70°C for 4 h. A
- 10 further portion of NaH (29 mg 60% dispersion in mineral oil) followed by 2-chloro-5-bromopyrimidine (138 mg) was added. Heating and stirring was continued for 25 h. Eventually, the reaction mixture was cooled to rt, diluted with EA (75 ml) and washed with 10% aqueous citric acid solution (50 ml) followed by water (50 ml). The organic layer was evaporated and the remaining residue was purified on
- 15 prep. tlc plates (silica gel, 0.5 mm layer thickness) developing with EA:methanol:sat. aqueous ammonia 8:2:1. This furnished 2-phenylethanesulfonic acid {5-(2-methoxy-phenoxy)-6-[2-(5-bromopyrimidin-2-yloxy)-ethoxy]-2-(2-pyrimidinyl)-4-pyrimidinyl]-amide as a pale yellow powder. LC-MS:  $t_R = 5.11$  min,  $[M+1]^+ = 680.23$ ,  $[M-1]^- = 678.36$ .

20

**Example 6**

To a suspension of NaH (29 mg, 60% dispersion in mineral oil) in a mixture of DMF and THF (2.5 ml each) was added 2-phenylethanesulfonic acid [6-(2-hydroxy-ethoxy)-5-(2-methoxy-phenoxy)-2-(2-pyrimidinyl)-4-pyrimidinyl]-amide (150 mg). After gas-evolution had ceased, 4,6-dimethoxy-2-methylsulfonylpyrimidine (156 mg) was added. The resulting yellow suspension was stirred at 70°C for 4 h. Eventually, the reaction mixture was cooled to rt, diluted with EA (75 ml) and washed with 10% aqueous citric acid solution (50 ml) followed by water (50 ml). The organic layer was evaporated and the remaining residue was purified on prep. tlc plates (silica gel, 0.5 mm layer thickness) developing with EA:methanol:sat. aqueous ammonia 8:2:1. This furnished 2-phenylethanesulfonic acid {5-(2-methoxy-phenoxy)-6-[2-(4,6-dimethoxy-pyrimidin-2-yloxy)-ethoxy]-2-(2-pyrimidinyl)-4-pyrimidinyl}-amide as a pale yellow powder. LC-MS:  $t_R = 5.21$  min,  $[M+1]^+ = 662.69$ ,  $[M-1]^- = 660.23$ .

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#### Example 7 - 86

The Examples represented in Table 1 to Table 8 have been prepared in analogy to the Examples 1 to 6 using the Referential Examples 1 to 29 as starting materials.

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Table 1:

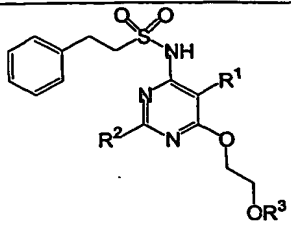
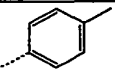
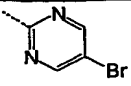
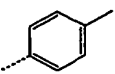
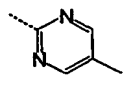
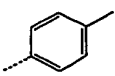

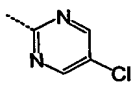
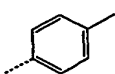

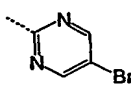
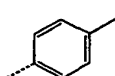

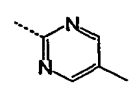
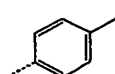

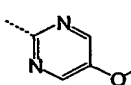
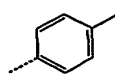

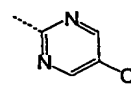
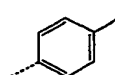

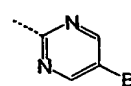
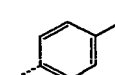

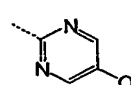
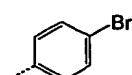
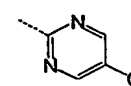
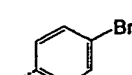
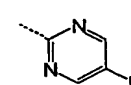
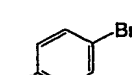
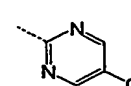
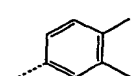
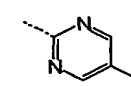
					
Ex. No	Ref. Ex. No	R <sup>1</sup>	R <sup>2</sup>	R <sup>3</sup>	LC-MS
7	1		H		t <sub>R</sub> = 5.41 min [M+H] <sup>+</sup> : 570.22 [M-H] <sup>-</sup> : 568.16
8	1		H		t <sub>R</sub> = 5.11 min [M+H] <sup>+</sup> : - [M-H] <sup>-</sup> : 504.11
9	2				t <sub>R</sub> = 5.26 min [M+H] <sup>+</sup> : 530.32 [M-H] <sup>-</sup> : -
10	2				t <sub>R</sub> = 5.21 min [M+H] <sup>+</sup> : 647.10 [M-H] <sup>-</sup> : 646.54
11	2				t <sub>R</sub> = 4.84 min [M+H] <sup>+</sup> : 583.22 [M-H] <sup>-</sup> : 581.18
12	2				t <sub>R</sub> = 4.82 min [M+H] <sup>+</sup> : 599.34 [M-H] <sup>-</sup> : 597.45
13	3				t <sub>R</sub> = 5.28 min [M+H] <sup>+</sup> : 604.11 [M-H] <sup>-</sup> : 602.42
14	3				t <sub>R</sub> = 5.33 min [M+H] <sup>+</sup> : 647.96 [M-H+2] <sup>-</sup> : 647.52
15	3				t <sub>R</sub> = 4.97 min [M+H] <sup>+</sup> : 600.19 [M-H] <sup>-</sup> : 598.28
16	4		H		t <sub>R</sub> = 5.51 min [M+H] <sup>+</sup> : 589.97 [M-H] <sup>-</sup> : 587.47
17	4		H		t <sub>R</sub> = 5.56 min [M+2+H] <sup>+</sup> : 636.23 [M-H] <sup>-</sup> : 632.12
18	4		H		t <sub>R</sub> = 5.18 min [M+H+2] <sup>+</sup> : 587.88 [M-H] <sup>-</sup> : 583.91
19	5		H		t <sub>R</sub> = 5.59 min [M+H] <sup>+</sup> : 584.10 [M-H+2] <sup>-</sup> : 583.54

Table 1: continued

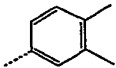
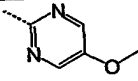
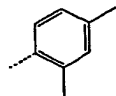
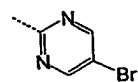
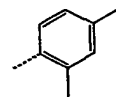
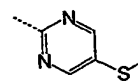
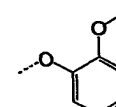


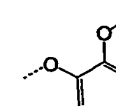

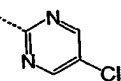
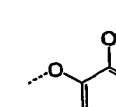

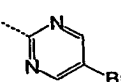
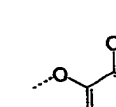

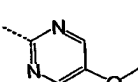
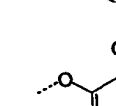

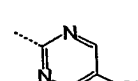
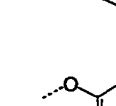

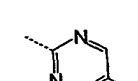
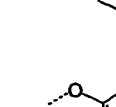

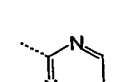


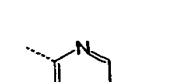
Ex. No	Ref. Ex. No	R <sup>1</sup>	R <sup>2</sup>	R <sup>3</sup>	LC-MS
20	5		H		t <sub>R</sub> = 5.24 min [M+H] <sup>+</sup> : 536.32 [M-H] <sup>-</sup> : 534.40
21	6		H		t <sub>R</sub> = 5.55 min [M+H] <sup>+</sup> : 584.18 [M-H] <sup>-</sup> : 582.12
22	6		H		t <sub>R</sub> = 5.46 min [M+H] <sup>+</sup> : 552.03 [M-H] <sup>-</sup> : 550.15
23	7				t <sub>R</sub> = 4.47 min [M+H] <sup>+</sup> : 600.95 [M-H] <sup>-</sup> : 598.84
24	7				t <sub>R</sub> = 4.97 min [M+H] <sup>+</sup> : 635.16 [M-H] <sup>-</sup> : 632.99
25	7				t <sub>R</sub> = 5.02 min [M+H] <sup>+</sup> : 679.17 [M-H] <sup>-</sup> : -
26	7				t <sub>R</sub> = 4.68 min [M+H] <sup>+</sup> : 631.30 [M-H] <sup>-</sup> : 629.22
27	8				t <sub>R</sub> = 4.90 min [M+H] <sup>+</sup> : 636.17 [M-H] <sup>-</sup> : 634.43
28	8				t <sub>R</sub> = 4.73 min [M+H] <sup>+</sup> : 614.10 [M-H] <sup>-</sup> : 612.15
29	8				t <sub>R</sub> = 4.68 min [M+H] <sup>+</sup> : 632.33 [M-H] <sup>-</sup> : 630.00
30	8				t <sub>R</sub> = 5.08 min [M+H] <sup>+</sup> : 670.19 [M-H] <sup>-</sup> : 668.34

Table 1: continued

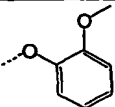

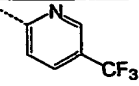
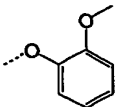


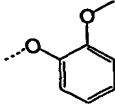

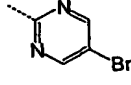
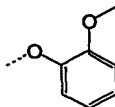

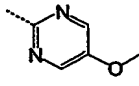
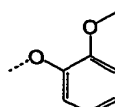

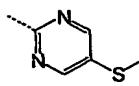
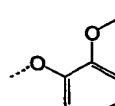


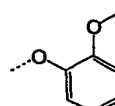

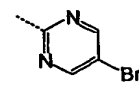
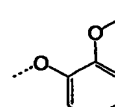

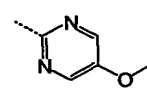
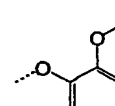

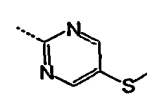
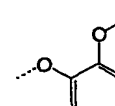

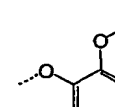
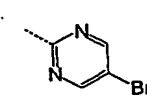
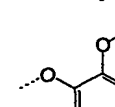
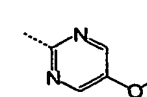
Ex. No	Ref. Ex. No	R <sup>1</sup>	R <sup>2</sup>	R <sup>3</sup>	LC-MS
31	8				$t_R = 5.42$ min [M+H] <sup>+</sup> : 669.05 [M-H] <sup>-</sup> : 666.80
32	9				$t_R = 4.70$ min [M+H] <sup>+</sup> : 602.25 [M-H] <sup>-</sup> : 600.18
33	9				$t_R = 5.19$ min [M+H+2] <sup>+</sup> : 681.96 [M-H] <sup>-</sup> : 678.21
34	9				$t_R = 4.88$ min [M+H] <sup>+</sup> : 632.24 [M-H] <sup>-</sup> : 630.35
35	9				$t_R = 5.11$ min [M+H] <sup>+</sup> : 648.57 [M-H] <sup>-</sup> : 646.11
36	10				$t_R = 5.13$ min [M+H] <sup>+</sup> : 609.31 [M-H] <sup>-</sup> : 607.25
37	10				$t_R = 5.67$ min [M+H] <sup>+</sup> : 686.92 [M-H] <sup>-</sup> : 685.02
38	10				$t_R = 5.30$ min [M+H] <sup>+</sup> : 639.30 [M-H] <sup>-</sup> : 637.13
39	10				$t_R = 5.51$ min [M+H] <sup>+</sup> : 654.92 [M-H] <sup>-</sup> : 653.08
40	11		H		$t_R = 4.84$ min [M+Na] <sup>+</sup> : 524.32 [M-H] <sup>-</sup> : 522.00
41	11		H		$t_R = 5.38$ min [M+H] <sup>+</sup> : 601.95 [M-H] <sup>-</sup> : 600.33
42	11		H		$t_R = 5.01$ min [M+H] <sup>+</sup> : 554.14 [M-H] <sup>-</sup> : 552.03

Table 1: continued

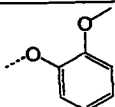
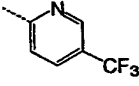
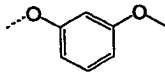

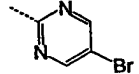
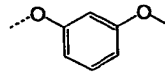

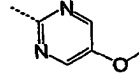
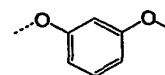
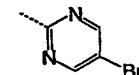
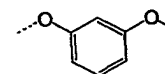
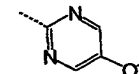
Ex. No	Ref. Ex. No	R <sup>1</sup>	R <sup>2</sup>	R <sup>3</sup>	LC-MS
43	11		H		$t_R = 5.75$ min [M+H] <sup>+</sup> : 591.44 [M-H] <sup>-</sup> : 588.99
44	12				$t_R = 5.52$ min [M+H] <sup>+</sup> : 687.24 [M-H] <sup>-</sup> : 685.44
45	12				$t_R = 5.18$ min [M+H] <sup>+</sup> : 639.32 [M-H] <sup>-</sup> : 637.14
46	13		H		$t_R = 5.25$ min [M+H] <sup>+</sup> : 602.00 [M-H] <sup>-</sup> : 599.99
47	13		H		$t_R = 4.89$ min [M+H] <sup>+</sup> : 553.98 [M-H] <sup>-</sup> : 551.97



Table 2:

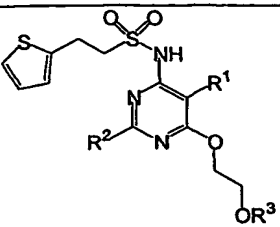
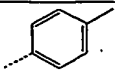
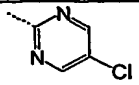
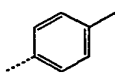
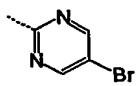
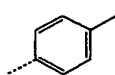
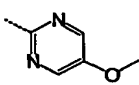
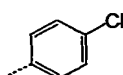
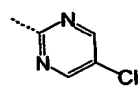
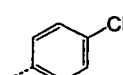
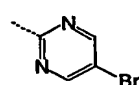
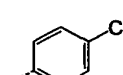
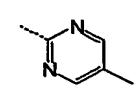
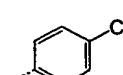
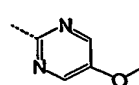
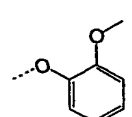

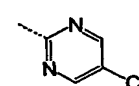
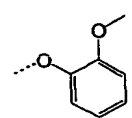

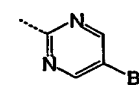
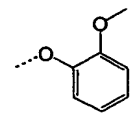

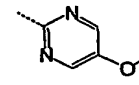
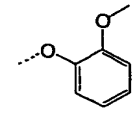
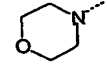
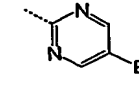
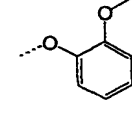
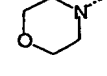
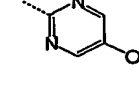
					
Ex. No	Ref. Ex. No	R <sup>1</sup>	R <sup>2</sup>	R <sup>3</sup>	LC-MS
48	14		H		t <sub>R</sub> = 5.24 min [M+H] <sup>+</sup> : 532.13 [M-H] <sup>-</sup> : 530.35
49	14		H		t <sub>R</sub> = 5.31 min [M+H] <sup>+</sup> : 576.07 [M-H] <sup>-</sup> : 574.19
50	14		H		t <sub>R</sub> = 4.95 min [M+H] <sup>+</sup> : 528.17 [M-H] <sup>-</sup> : 526.12
51	15		H		t <sub>R</sub> = 5.28 min [M+H] <sup>+</sup> : 552.39 [M-H] <sup>-</sup> : 550.27
52	15		H		t <sub>R</sub> = 5.33 min [M+H+2] <sup>+</sup> : 598.29 [M-H] <sup>-</sup> : 595.43
53	15		H		t <sub>R</sub> = 5.07 min [M+H] <sup>+</sup> : 532.29 [M-H] <sup>-</sup> : 529.95
54	15		H		t <sub>R</sub> = 4.99 min [M+H] <sup>+</sup> : 547.89 [M-H] <sup>-</sup> : 545.91
55	16				t <sub>R</sub> = 5.07 min [M+H] <sup>+</sup> : 641.16 [M-H] <sup>-</sup> : 638.75
56	16				t <sub>R</sub> = 4.96 min [M+H] <sup>+</sup> : 685.12 [M-H] <sup>-</sup> : 683.33
57	16				t <sub>R</sub> = 4.62 min [M+H] <sup>+</sup> : 637.18 [M-H] <sup>-</sup> : 635.04
58	17				t <sub>R</sub> = 5.56 min [M+H] <sup>+</sup> : 693.12 [M-H] <sup>-</sup> : 691.31
59	17				t <sub>R</sub> = 5.21 min [M+H] <sup>+</sup> : 645.35 [M-H] <sup>-</sup> : 643.48

Table 3:

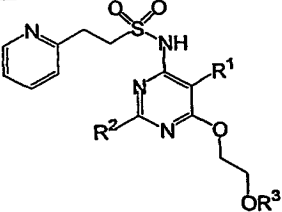
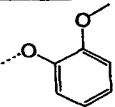


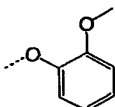

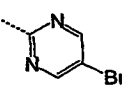
					
Ex. No	Ref. Ex. No	R <sup>1</sup>	R <sup>2</sup>	R <sup>3</sup>	LC-MS
60	18				$t_R = 3.56 \text{ min}$ [M+H] <sup>+</sup> : 602.40 [M-H] <sup>-</sup> : 600.14
61	18				$t_R = 4.14 \text{ min}$ [M+H+2] <sup>+</sup> : 682.26 [M-H] <sup>-</sup> : 678.23

Table 4:

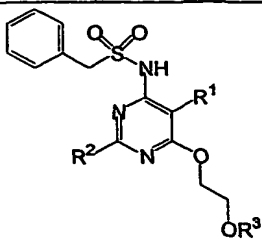
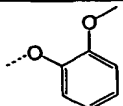


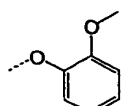

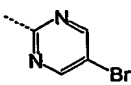
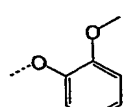

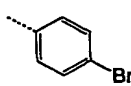
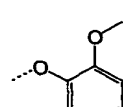


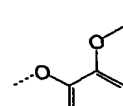

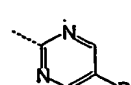
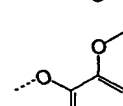

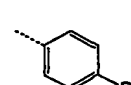
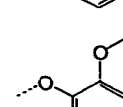


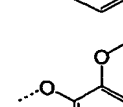

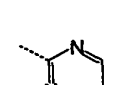
					
Ex. No	Ref. Ex. No	R <sup>1</sup>	R <sup>2</sup>	R <sup>3</sup>	LC-MS
62	19				$t_R = 4.46$ min [M+H] <sup>+</sup> : 587.43 [M-H] <sup>-</sup> : 585.40
63	19				$t_R = 5.00$ min [M+H+2] <sup>+</sup> : 667.12 [M-H+2] <sup>-</sup> : 665.47
64	19				$t_R = 5.71$ min [M+H+2] <sup>+</sup> : 665.39 [M-H+2] <sup>-</sup> : 662.71
65	20				$t_R = 4.43$ min [M+H] <sup>+</sup> : 588.66 [M-H] <sup>-</sup> : 586.33
66	20				$t_R = 4.93$ min [M+H+2] <sup>+</sup> : 668.21 [M-H+2] <sup>-</sup> : 665.81
67	20				$t_R = 5.52$ min [M+H+2] <sup>+</sup> : 666.29 [M-H] <sup>-</sup> : 662.25
68	21				$t_R = 5.01$ min [M+H] <sup>+</sup> : 595.27 [M-H] <sup>-</sup> : 593.18
69	21				$t_R = 5.46$ min [M+H] <sup>+</sup> : 673.11 [M-H+2] <sup>-</sup> : 673.42

Table 5:

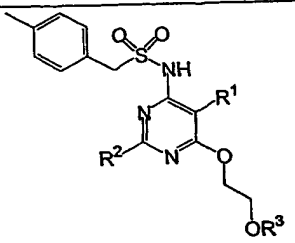
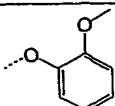


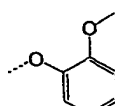

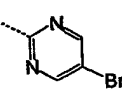
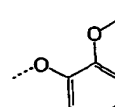


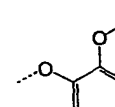

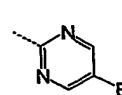
					
Ex. No	Ref. Ex. No	R <sup>1</sup>	R <sup>2</sup>	R <sup>3</sup>	LC-MS
70	22				$t_R = 4.58 \text{ min}$ $[M+H]^+ : 601.39$ $[M-H]^- : 599.12$
71	22				$t_R = 5.28 \text{ min}$ $[M+H+2]^+ : 681.24$ $[M-H]^- : 677.00$
72	23				$t_R = 4.76 \text{ min}$ $[M+H]^+ : 602.20$ $[M-H]^- : 600.18$
73	23				$t_R = 5.24 \text{ min}$ $[M+H]^+ : 680.13$ $[M-H]^- : 678.14$

Table 6

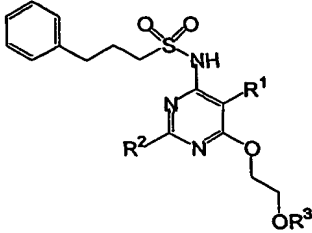
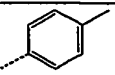

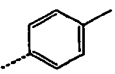
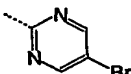
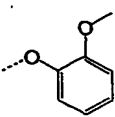


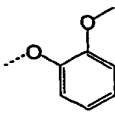

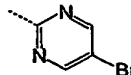
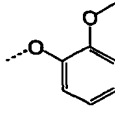


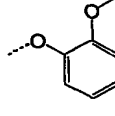

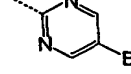
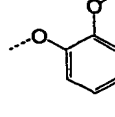


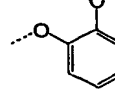

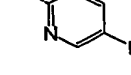
					
Ex. No	Ref. Ex. No	R <sup>1</sup>	R <sup>2</sup>	R <sup>3</sup>	LC-MS
74	24		H		$t_R = 5.13$ min [M+H] <sup>+</sup> : 506.25 [M-H] <sup>-</sup> : 504.26
75	24		H		$t_R = 5.62$ min [M+H+2] <sup>+</sup> : 586.33 [M-H] <sup>-</sup> : 582.11
76	25				$t_R = 4.56$ min [M+H] <sup>+</sup> : 615.53 [M-H] <sup>-</sup> : 613.40
77	25				$t_R = 5.09$ min [M+H] <sup>+</sup> : 693.22 [M-H] <sup>-</sup> : 691.07
78	26				$t_R = 4.58$ min [M+H] <sup>+</sup> : 616.54 [M-H] <sup>-</sup> : 614.29
79	26				$t_R = 5.09$ min [M+H] <sup>+</sup> : 694.42 [M-H] <sup>-</sup> : 692.21
80	27				$t_R = 5.20$ min [M+H] <sup>+</sup> : 623.37 [M-H] <sup>-</sup> : 621.19
81	27				$t_R = 5.66$ min [M+H+2] <sup>+</sup> : 703.51 [M-H+2] <sup>-</sup> : 701.52

Table 7:

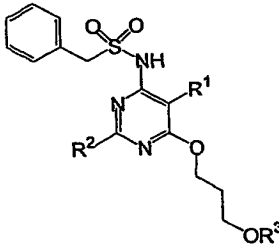
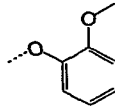


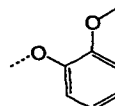

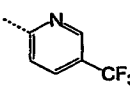
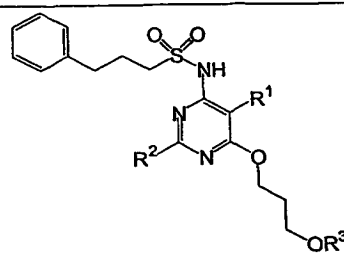
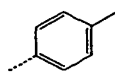
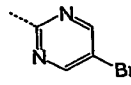
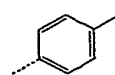
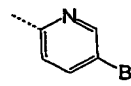
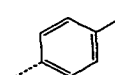
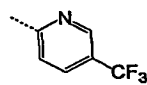
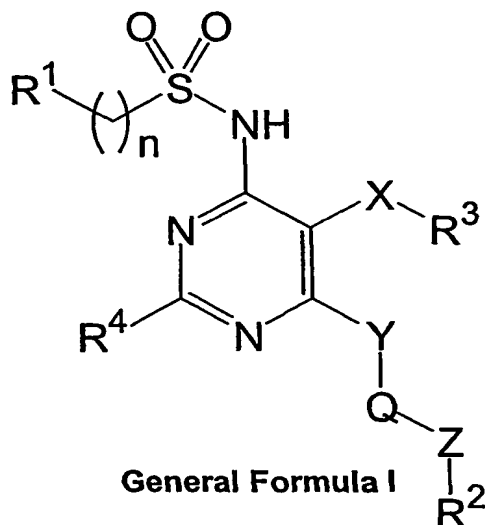
					
Ex. No	Ref. Ex. No	R <sup>1</sup>	R <sup>2</sup>	R <sup>3</sup>	LC-MS
82	28				t <sub>R</sub> = 4.55 min [M+H] <sup>+</sup> : 602.43 [M-H] <sup>-</sup> : 600.26
83	28				t <sub>R</sub> = 5.59 min [M+H] <sup>+</sup> : 669.49 [M-H] <sup>-</sup> : 667.40

Table 8:

					
Ex. No	Ref. Ex. No	R <sup>1</sup>	R <sup>2</sup>	R <sup>3</sup>	LC-MS
84	29		H		t <sub>R</sub> = 5.71 min [M+H+2] <sup>+</sup> : 600.18 [M-H] <sup>-</sup> : 596.05
85	29		H		t <sub>R</sub> = 6.21 min [M+H+2] <sup>+</sup> : 598.96 [M-H] <sup>-</sup> : 594.67
86	29		H		t <sub>R</sub> = 6.16 min [M+H] <sup>+</sup> : 587.19 [M-H] <sup>-</sup> : 584.74

## Claims

### 1. Compounds of the general formula I



5

wherein

$R^1$  and  $R^2$  represent aryl; heteroaryl;

$R^3$  represents phenyl; mono-, di- or tri-substituted phenyl substituted with lower alkyl, lower alkenyl, lower alkynyl, phenyl, lower alkoxy, amino, lower alkylamino, amino-lower alkyl, trifluoromethyl, trifluoromethoxy, halogen, lower alkylthio, hydroxy, hydroxy-lower alkyl, cyano, carboxyl, lower alkanoyl, formyl; benzofuranyl; aryl; heteroaryl;

$R^4$  represents hydrogen; halogen; trifluoromethyl; lower alkyl; lower alkyl-amino; lower alkoxy; lower alkyl-sulfonyl; lower alkyl-sulfinyl; lower alkylthio; lower alkylthio-lower alkyl; hydroxy-lower alkyl; lower alkyl-oxy-lower alkyl; hydroxy-lower alkyl-oxy-lower alkyl; hydroxy-lower alkyl-amino; lower alkyl-amino-lower alkyl; amino; di-lower alkyl-amino; [N-(hydroxy-lower alkyl)-N-(lower alkyl)]-amino; aryl; aryl-amino; aryl-lower alkyl-amino; aryl-thio; aryl-lower alkyl-thio; aryloxy; aryl-lower alkyl-oxy; aryl-lower alkyl; aryl-sulfinyl; heteroaryl; heteroaryl-oxy; heteroaryl-lower alkyl-oxy; heteroaryl-amino; heteroaryl-lower alkyl-amino; heteroaryl-thio; heteroaryl-lower alkyl-thio; heteroaryl-lower alkyl; heteroaryl-sulfinyl; heterocyclyl; heterocyclyl-lower alkyl-oxy; heterocyclyl-oxy; heterocyclyl-

amino; heterocyclyl-lower alkyl-amino; heterocyclyl-thio; heterocyclyl-lower alkyl-thio; heterocyclyl-lower alkyl; heterocyclyl-sulfinyl; cycloalkyl; cycloalkyl-oxy; cycloalkyl-lower alkyl-oxy; cycloalkyl-amino; cycloalkyl-lower alkyl-amino; cycloalkyl-thio; cycloalkyl-lower alkyl-thio; cycloalkyl-lower alkyl; cycloalkyl-sulfinyl;

X represents oxygen; sulfur; NH; CH<sub>2</sub> or a bond;

Y represents oxygen; sulfur or -NH-;

Z represents oxygen; sulfur, -NH- or a bond;

Q represents -(CH<sub>2</sub>)<sub>k</sub>-; -(CH<sub>2</sub>)<sub>m</sub>-C≡C-(CH<sub>2</sub>)<sub>p</sub>-, in case p represents 0 (zero), Z represents a bond; -CH<sub>2</sub>-cyclopropylen-CH<sub>2</sub>-;

k represents the numbers 2, 3, 4, 5, or 6;

m represents the numbers 1, 2, or 3;

p represents the numbers 0, 1, 2 or 3;

n represents the numbers 1, 2, or 3;

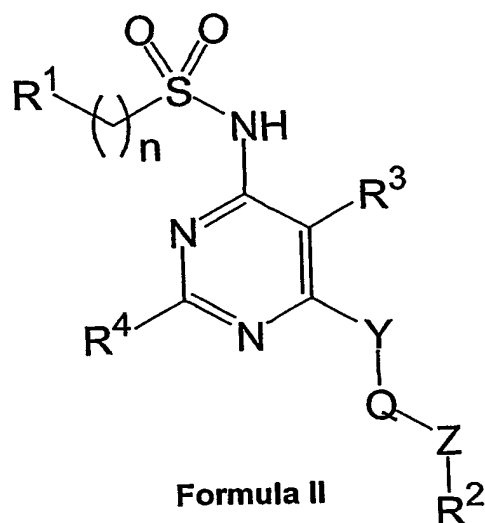
and pure diastereomers, mixtures of diastereomers, diastereomeric racemates, mixtures of diastereomeric racemates and the meso-forms and pharmaceutically acceptable salts thereof.

2. Compounds of the formula I in claim 1, wherein R<sup>1</sup>, R<sup>2</sup>, R<sup>4</sup>, Q, Y, Z and n are as defined in general formula I in claim 1, X, represents oxygen and R<sup>3</sup> represents phenyl or mono-substituted phenyl substituted with halogen, lower alkyl, lower alkenyl, lower alkoxy, amino, lower alkyl-amino, lower alkyl-thio, hydroxy, hydroxymethyl and lower alkanoyl; and pharmaceutically acceptable salts of compounds of general formula I.



3. Compounds of the formula I in claim 1, wherein  $R^1$ ,  $R^2$ ,  $R^4$ ,  $Q$ , and  $n$  are as defined in general formula I in claim 1,  $X$ ,  $Y$  and  $Z$  represent oxygen and  $R^3$  represents phenyl or mono-substituted phenyl substituted with halogen, lower alkyl, lower alkenyl, lower alkoxy, amino, lower alkyl-amino, lower alkyl-thio, hydroxy, hydroxymethyl and lower alkanoyl;  
5 and pharmaceutically acceptable salts of compounds of general formula I.
4. Compounds of the general formula I in claim 1, wherein  $R^1$ ,  $R^2$ ,  $R^4$ , and  $n$  are as defined in general formula I in claim 1,  $X$ ,  $Y$  and  $Z$  represent oxygen,  $Q$  represents  $-(CH_2)_k-$  with  $k = 2$  or  $3$  and  $R^3$  represents phenyl or mono-substituted phenyl substituted with halogen, lower alkyl, lower alkenyl, lower alkoxy, amino, lower alkyl-amino, lower alkyl-thio, hydroxy, hydroxymethyl and lower alkanoyl;  
10 and pharmaceutically acceptable salts of compounds of general formula I.
5. Compounds of the general formula I in claim 1, wherein  $R^1$ ,  $R^4$ , and  $n$  are as defined in general formula I in claim 1,  $X$ ,  $Y$  and  $Z$  represent oxygen,  $Q$  represents  $-(CH_2)_k-$  with  $k = 2$  or  $3$ ,  $R^2$  represents heteroaryl and  $R^3$  represents phenyl or mono-substituted phenyl substituted with halogen, lower alkyl, lower alkenyl, lower alkoxy, amino, lower alkyl-amino, lower alkyl-thio, hydroxy, hydroxymethyl and lower alkanoyl;  
15 20 and pharmaceutically acceptable salts of compounds of general formula I.
6. Compounds of the general formula I in claim 1, wherein  $R^1$ ,  $R^4$ , and  $n$  are as defined in general formula I in claim 1,  $X$ ,  $Y$  and  $Z$  represent oxygen,  $Q$  represents  $-(CH_2)_2-$ ,  $R^2$  represents heteroaryl,  $R^3$  represents phenyl or mono-substituted phenyl substituted with halogen, lower alkyl, lower alkenyl, methoxy, amino, lower alkyl-amino, lower alkyl-thio, hydroxy, hydroxymethyl and lower alkanoyl;  
25 and pharmaceutically acceptable salts of compounds of general formula I.

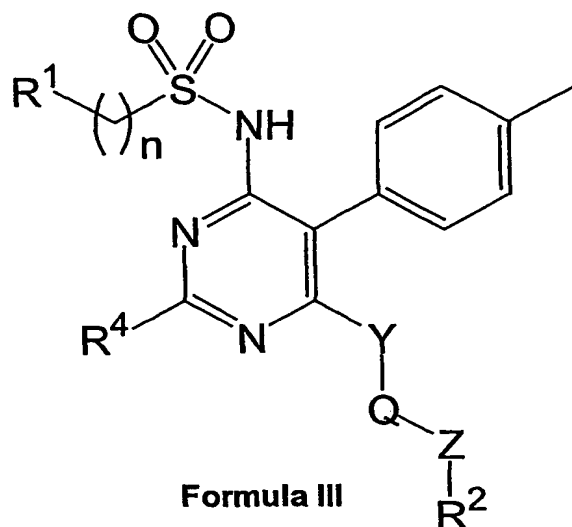
## 7. Compounds of formula II



5

wherein  $R^1$ ,  $R^2$ ,  $R^3$ ,  $R^4$ , Y, Q, Z and n are as defined in general formula I in claim 1 above,  
and pharmaceutically acceptable salts of compounds of formula II.

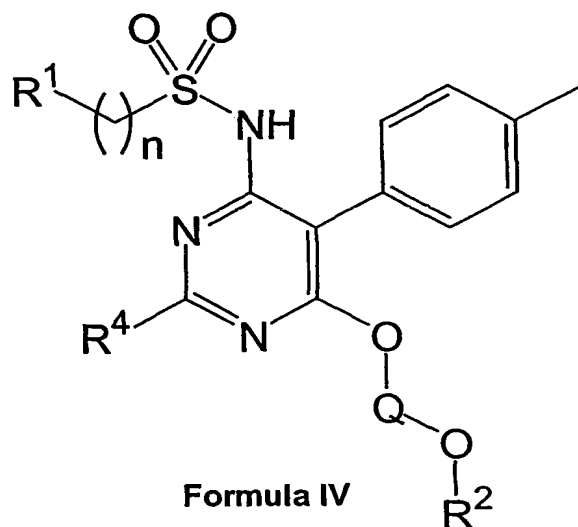
## 8. Compounds of formula III



- 5 wherein  $R^1$ ,  $R^2$ ,  $R^4$ ,  $Y$ ,  $Q$ ,  $Z$  and  $n$  are as defined in general formula I in claim 1 above,  
and pharmaceutically acceptable salts of compounds of formula III.

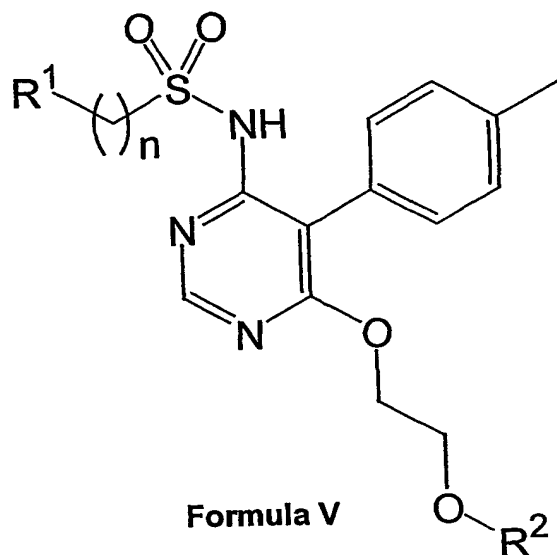
## 9. Compounds of the formula IV

10



- wherein  $R^1$ ,  $R^2$ ,  $R^4$ ,  $Q$  and  $n$  are as defined in general formula I in claim 1 above,  
and pharmaceutically acceptable salts of compounds of formula IV.

## 10. Compounds of formula V



5

wherein  $R^1$ ,  $R^2$  and  $n$  are as defined in general formula I in claim 1 above,  
and pharmaceutically acceptable salts thereof.

11. Compounds according to claim 10, wherein  $R^2$  in formula V represents  
10 heteroaryl and pharmaceutically acceptable salts thereof.

12. Compounds according to any one of claims 1 to 11 selected from the group  
consisting of

- 15        2-Phenyl-ethanesulfonic acid {6-[2-(5-bromo-pyrimidin-2-yloxy)-ethoxy]-5-  
p-tolyl-pyrimidin-4-yl}-amide,  
          2-Phenyl-ethanesulfonic acid {6-[2-(5-bromo-pyrimidin-2-yloxy)-ethoxy]-5-  
p-tolyl-[2,2']bipyrimidinyl-4-yl}-amide,  
          2-Phenyl-ethanesulfonic acid {5-(2-methoxy-phenoxy)-6-[2-(pyrimidin-2-  
20 yloxy)-ethoxy]-[2,2']bipyrimidinyl-4-yl}-amide,  
          2-Phenyl-ethanesulfonic acid {5-(2-methoxy-phenoxy)-6-[2-(5-methoxy-  
pyrimidin-2-yloxy)-ethoxy]-[2,2']bipyrimidinyl-4-yl}-amide,

2-Thiophen-2-yl-ethanesulfonic acid [6-[2-(5-bromo-pyrimidin-2-yloxy)-ethoxy]-5-(4-chloro-phenyl)-pyrimidin-4-yl]-amide,

2-Thiophen-2-yl-ethanesulfonic acid {5-(4-chloro-phenyl)-6-[2-(5-methoxy-pyrimidin-2-yloxy)-ethoxy]-pyrimidin-4-yl}-amide,

5 2-Pyridin-2-yl-ethanesulfonic acid [6-[2-(5-bromo-pyrimidin-2-yloxy)-ethoxy]-5-(2-methoxy-phenoxy)-2-pyridin-4-yl-pyrimidin-4-yl]-amide,

and pharmaceutically acceptable salts thereof.

10

13. A compound as described as final product in any of the Examples 1-86

14. Pharmaceutical compositions for the treatment of disorders which are associated with a role of endothelin, especially circulatory disorders such as  
15 hypertension, ischaemia, vasospasm and angina pectoris and proliferative disorders such as cancer, containing a compound of any one of claims 1 to 13 and usual carrier materials and adjuvants.

15. Pharmaceutical compositions for the treatment of disorders which are  
20 associated with a role of endothelin, such as migraine, asthma or inflammatory disorders, containing a compound of any one of claims 1 to 13 and usual carrier materials and adjuvants.

16. The compounds of any one of the claims 1 to 13 for use as medicaments for  
25 the treatment of disorders which are associated with a role of endothelin, especially circulatory disorders such as hypertension, ischaemia, vasospasm and angina pectoris, proliferative disorders such as cancer, migraine and inflammatory disorders.

30 17. The compounds of any one of the claims 1 to 13 for use as medicaments for the treatment of disorders which are associated with a role of endothelin, and require mixed ET<sub>A</sub> and ET<sub>B</sub> blocking for treatment.

18. The compounds of any one of the claims 1 to 13 for use as medicaments for the treatment of disorders which are associated with a role of endothelin, and require selective ET<sub>A</sub> blocking for treatment.
- 5 19. The compounds of any one of the claims 1 to 13 for use as medicaments for the treatment of disorders which are associated with a role of endothelin, and require selective ET<sub>B</sub> blocking for treatment.
- 10 20. The use of one or more compounds of any one of claims 1 to 13 as active ingredients for the production of pharmaceutical compositions for the treatment of disorders associated with a role of endothelin, especially circulatory disorders such as hypertension, ischaemia, vasospasm and angina pectoris and proliferative disorders such as cancer.
- 15 21. The use of one or more compounds of any one of claims 1 to 13 as active ingredients for the production of pharmaceutical compositions for the treatment of disorders associated with endothelin activities, such as migraine, asthma or inflammatory disorders.
- 20 22. A process for the manufacture of pharmaceutical compositions for the treatment of disorders associated with a role of endothelin containing one or more compounds as claimed in any one of claims 1 to 13 as active ingredients which process comprises mixing one or more active ingredient with pharmaceutically acceptable excipients in a manner known per se.
- 25 23. The invention as hereinbefore described.

# INTERNATIONAL SEARCH REPORT

International Application No  
PCT/EP 01/09894

<b>A. CLASSIFICATION OF SUBJECT MATTER</b> IPC 7 C07D239/46 A61K31/505 A61P9/10 C07D409/14		
According to International Patent Classification (IPC) or to both national classification and IPC		
<b>B. FIELDS SEARCHED</b> Minimum documentation searched (classification system followed by classification symbols) IPC 7 C07D		
Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched		
Electronic data base consulted during the international search (name of data base and, where practical, search terms used) EPO-Internal, CHEM ABS Data		
<b>C. DOCUMENTS CONSIDERED TO BE RELEVANT</b>		
Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	EP 0 743 307 A (TANABE SEIYAKU CO) 20 November 1996 (1996-11-20) cited in the application page 30; example 5; table 1 page 32; example 9	1-12, 14-22
Y	EP 0 882 719 A (YAMANOUCHI PHARMA CO LTD) 9 December 1998 (1998-12-09) cited in the application page 31; example 1; table 5	1-12, 14-22
Y	WO 98 57938 A (FUJIMORI AKIRA; HARADA HIRONORI (JP); KAZAMI JUN ICHI (JP); WATANU) 23 December 1998 (1998-12-23) page 19; examples 2,4; table 3	1-12, 14-22
<input type="checkbox"/> Further documents are listed in the continuation of box C. <span style="margin-left: 100px;"><input checked="" type="checkbox"/> Patent family members are listed in annex.</span>		
<div style="display: flex; justify-content: space-between;"> <div style="width: 45%;"> <p>* Special categories of cited documents :</p> <p>*A* document defining the general state of the art which is not considered to be of particular relevance</p> <p>*E* earlier document but published on or after the international filing date</p> <p>*L* document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)</p> <p>*O* document referring to an oral disclosure, use, exhibition or other means</p> <p>*P* document published prior to the international filing date but later than the priority date claimed</p> </div> <div style="width: 45%;"> <p>*T* later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention</p> <p>*X* document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone</p> <p>*Y* document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.</p> <p>*&amp;* document member of the same patent family</p> </div> </div>		
Date of the actual completion of the international search  <div style="text-align: center; font-size: 1.2em;">26 February 2002</div>		Date of mailing of the international search report  <div style="text-align: center; font-size: 1.2em;">11/03/2002</div>
Name and mailing address of the ISA European Patent Office, P.B. 5818 Patentlaan 2 NL - 2280 HV Rijswijk Tel. (+31-70) 340-2040, Tx. 31 651 epo nl, Fax: (+31-70) 340-3016		Authorized officer  <div style="text-align: center; font-size: 1.2em;">Timmermans, M</div>

## INTERNATIONAL SEARCH REPORT

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Patent document cited in search report		Publication date	Patent family member(s)	Publication date
EP 0743307	A	20-11-1996	JP 3013752 B2	28-02-2000
			JP 8311043 A	26-11-1996
			AT 205482 T	15-09-2001
			CA 2176776 A1	17-11-1996
			CN 1146990 A	09-04-1997
			DE 69615085 D1	18-10-2001
			DK 743307 T3	26-11-2001
			EP 0743307 A1	20-11-1996
			EP 0959072 A1	24-11-1999
			EP 0959073 A1	24-11-1999
			PT 743307 T	28-12-2001
			SG 47141 A1	20-03-1998
			US 5739333 A	14-04-1998
			JP 3013778 B2	28-02-2000
			JP 9059261 A	04-03-1997
EP 0882719	A	09-12-1998	AT 201202 T	15-06-2001
			AU 703386 B2	25-03-1999
			AU 1171097 A	14-07-1997
			BR 9612061 A	23-02-1999
			DE 69612874 D1	21-06-2001
			DE 69612874 T2	04-10-2001
			EP 0882719 A1	09-12-1998
			JP 3087968 B2	18-09-2000
			US 6083955 A	04-07-2000
			CA 2238723 A1	26-06-1997
			ES 2156305 T3	16-06-2001
			HU 0000474 A2	28-12-2000
			WO 9722595 A1	26-06-1997
			TW 414792 B	11-12-2000
WO 9857938	A	23-12-1998	AU 8035698 A	04-01-1999
			WO 9857938 A1	23-12-1998



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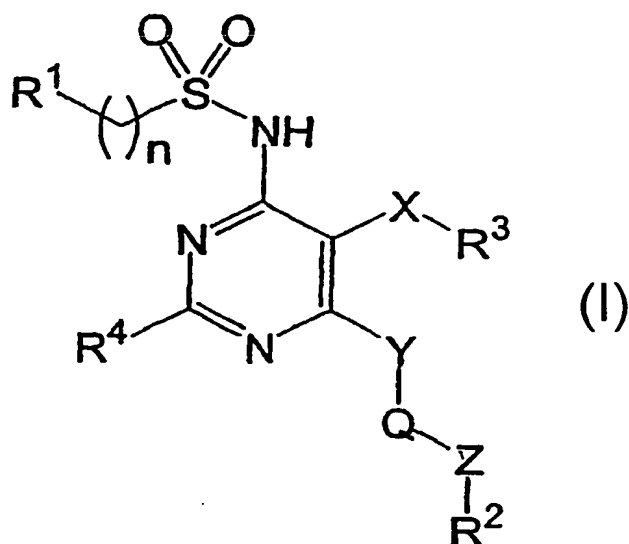
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(54) Title: ARYLALKANE-SULFONAMIDES HAVING ENDOTHELIN-ANTAGONIST ACTIVITY



(57) Abstract: The invention relates to novel aryl-alkane-sulfonamides of the general formula (I) and their use as active ingredients in the preparation of pharmaceutical compositions. The invention also concerns related aspects including processes for the preparation of the compounds, pharmaceutical compositions containing one or more of those compounds and especially their use as endothelin receptor antagonists.

WO 02/24665 A1

37X2

WO 02/24665 A1



**(15) Information about Correction:**  
see PCT Gazette No. 23/2002 of 6 June 2002, Section II

*For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.*